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RESULT 7
SS1364
sperm tail-specific protein msc101(2) - fruit fly (*Drosophila hydei*)
C/Species: *Drosophila hydei*
C/Date: 19-Jul-1996 #sequence_revision 26-Jul-1996 #text_change 17-Mar-1999
C/Accession: SS1364; S34154
R./Neesen, J.; Padmanabhan, S.; Buenemann, H.
Eur. J. Biochem. 225, 1089-1095, 1994
A./Title: Randomly arranged repeats of a novel highly charged 16-amino-acid motif represses
alpha-helical rods within the extremely elongated spermatzoa of *Drosophila hydei*.
A./Reference number: SS1364; MUID:95045538; PMID:7957199
A/Accession: SS1364
A./Status: nucleic acid sequence not shown
A./Molecule type: DNA
A./Residues: 1-1390 <NEE>
A./Cross-references: EMBL:X73481
R./Neesen, J.; Heinlein, U.A.O.; Buenemann, H.
submitted to the EMBL Data Library, June 1993
A./Reference number: S34153
A/Accession: S34154
A./Molecule type: DNA
A./Residues: 1-163, 'E', 164-236, 'Q', 237-254, 257-320, 'E', 321-1390 <NEW>
A./Cross-references: EMBL:X73481; NID:g313201; PID:g313202
C/Genetics:
A./Gene: msc101(2)
A./Cross-references: FlyBase:FBgn0011816

	Query Match	42.3%	Score 90;	DB 2;	Length 1390;
	Best Local Similarity	57.8%	Pred No. 0.41;		
	Matches	26;	Conservative	4;	Mismatches 11; Indels 4; Gaps 2.
Oy	2 KCTAKGAAEK- AKKAYDALEAK--AAKTCKAALAEQAANKEA	42			
	:::::::::::::::::::::	:			
b	662 KETKEKKCEKAARKKEAEKKCAEAKKEKEKAEEKKCEEA	706			
	:::::::::::::::::::::	:			

RESULT 8
A28100
histone H1-beta, embryonic - sea urchin (Strongylocentrotus purpuratus)
C:Species: Strongylocentrotus purpuratus (purple urchin)
C:Date: 28-Aug-1989 #sequence_revision 28-Aug-1989 #text_change 23-Feb-1997
C:Accession: A28100
R:Lai, Z.C.; Childs, G.
Mol. Cell. Biol. 8, 1842-1844, 1988
A:Title: Characterization of the structure and transcriptional patterns of the gene encoding histone H1-beta in the sea urchin Strongylocentrotus purpuratus
A:Reference number: A28100; MUID:88246461; PMID:2837660
A:Accession: A28100
A:Molecule type: DNA
A:Residues: 1-211 <LAI>
A:Cross-references: GB:M20314
C:Superfamily: histone H1
;Keywords: chromosomal protein; DNA binding; embryo; nucleosome; nucleus

```

Query Match      42.0%; Score 89.5; DB 2, length 211,
Best Local Similarity 61.9%; Pred. No. 0.11;
Matches 26; Conservative 3; Mismatches 10; Indels 3; Gaps 2.

Oy      1 AKTAKKAKAEEKAKAYAAAEKKAKAEKAAAEKAAAEKAA 42
Db      139 SKTTTKKVKKKPAKKAKKAPKA-AKKAK--KPAKKAPPAKKAA 177

```

RESULT 9
G70673
probable hnpB - Mycobacterium tuberculosis (strain H37Rv)
C:Species: Mycobacterium tuberculosis
C:Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 20-Jun-2000
C:Accession: G70673
R:Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon, S.; Connor, R.; Davies, R.; Devlin, K.; Fellwell, T.; Gentile, S.; Hamlin, N.; Holroyd, S.; Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.
Nature 393, 537-544, 1998
A:Authors: Sgares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.

A:Title: Deciphering the biology of *Mycobacterium tuberculosis* from the complete genome
A:Reference number: A70500; MUID:98295987; PMID:9634230
A:Accession: G70673
A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-214 <COL>
A:Cross-references: GB:Z83018; GB:AL123456; NID:g3261671; PIDD:CAM05427.1; PID:g1694845
A:Experimental source: strain H37Rv
C:Genetics:
A:Gene: hupB
C:superfamily: histone H1

Query Match	42.0%	Score 89.5	DB 2	Length 214
Best Local Similarity	55.6%	Pred. No. 0.11		
Matches	25	Conservative	4	Mismatches 13
				Indels 3
				Gaps 1

```

RESULT 10
B87553
DNA topoisomerase I [imported] - Caulobacter crescentus
C|Species: Caulobacter crescentus
C|Date: 20-Apr-2001 #sequence_revision 20-Apr-2001 #text_change 10-May-2001
C|Accession: B87553
R|Nierman, W.C.; Feldblyum, T.V.; Paulsen, I.T.; Nelson, K.E.; Eisen, J.; Heidelberg, J.
B.; Laub, M.J.; DeBoy, R.T.; Dodson, R.J.; Durkin, A.S.; Gwinn, M.L.; Haft, D.H.; Kolon
n, J.; Esmolaev, M.; White, O.; Salzberg, S.L.; Shapiro, L.; Venter, J.C.; Fraser, C.M.
Proc. Natl. Acad. Sci. U.S.A. 98, 4116-4141, 2001
A|Title: Complete Genome Sequence of Caulobacter crescentus.
A|Reference number: A87249; MUID:21173698; PMID:11259647
A|Accession: B87553
A|Status: preliminary
A|Molecule type: DNA
A|Residues: 1899 <STO>
A|Cross-references: GB:AE005673; NID:gi342398; PIDD:AAK24422.1; GSPDB:GN00148
C|Genetic8:
A|Gene: CC2451
C|Superfamily: bacterial type I DNA topoisomerase

```

Query Match	41.3%	Score	88	DB	2	Length	899
Best Local Similarity	58.7%	Pred. No.	0.45				
Matches	27	Conservative	5	Mismatches	12	Indels	
						Gaps	2
QY	1	AKKAKAKAKAKAK-KAYKAAEAKKAKYEGQAQAAEKAAAKAAEAA	45				
DB	836	AKKAKAKAAAKKSKAKKESDAPAKKATA-KKPAKAKKPAKAKKAPKA	880				

RESULT 11
B43592
outer membrane protein TmpB - Treponema phagedenis
C|Species: Treponema phagedenis
C|Date: 30-Jan-1993 #sequence_revision 12-Mar-1993 #text_change 08-Oct-1999
C|Accession: B43592
R|Yelton, D. B.; Limberger, R. J.; Curci, K.; Malinovsky-Rummell, F.; Slivenski, L.; Schou
Infect. Immun. 59, 3685-3693, 1991
A|Title: Treponema phagedenis encodes and expresses homologs of the Treponema pallidum T
A|Reference number: B43592; NUID:91372983; PMID:1894368
A|Accession: B43592
A|Status: preliminary
A|Molecule type: DNA
A|Residues: 1-384 <YEL>
A|Cross-references: GB:M58563; NID:G15506; PIDN:AA27480.1; PID:G155067
A|Note: the authors translated the codon TTC for residue 316 as Tyr, and CGA for residue
C|Keywords: membrane protein

Query Match	40.6%	Score 86.5	DB 2	Length 384
Best Local Similarity	54.5%	Pred. No. 0.32		
Matches 24	Conservative	6	Mismatches 13	Indels 1
			Gaps	1

..! Lory, S.; Olson, M.V.

RESULT 2
TOL A ECOLI STANDARD; PRT; 421 AA.
ID TOL A ECOLI
AC P19934;
DT 01-FEB-1991 (Rel. 17, Created)
DT 01-FEB-1991 (Rel. 17, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE TOL A protein.
GN TOL A OR CIM OR EXCC OR LKY OR B0739.
OS Escherichia coli.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Escherichia.
OX NCBI_TaxId=562;
[1]
RP SEQUENCE FROM N.A.
RC STRAIN=JMI05;
RX MEDLINE=90078104; PubMed=2687247;
RA Levensgood S.K., Webster R.E.;
RT "Nucleotide sequences of the tolA and tolB genes and localization of
RT their products, components of a multistep translocation system in
RT Escherichia coli."
RL J. Bacteriol. 171:6600-6609(1989).
[2]
RP SEQUENCE FROM N.A.
RC STRAIN=K12 / MG1655;
RX MEDLINE=97426617; PubMed=9278503;
RA Blatner F.R., Plunkett G. III, Bloch C.A., Perna N.T., Burland V.,
RA Riley M., Collado-Vides J., Glasner J.D., Rode C.K., Mayhew G.F.,
RA Gregor J., Davis N.W., Kirkpatrick H.A., Goeden M.A., Rose D.J.,
RA Mau B., Shao Y.;
RT "The complete genome sequence of Escherichia coli K-12."
RL Science 277:1453-1474(1997).
[3]
RP SEQUENCE FROM N.A.
RC STRAIN=K12;
RX MEDLINE=97061202; PubMed=8905232;
RA Oshima T., Aida H., Baba T., Fujita K., Hayashi K., Honjo A.,
RA Ikemoto K., Inada T., Itoh T., Kajihara M., Kanai K., Kasahimoto K.,
RA Kimura S., Kitagawa M., Makino K., Masuda S., Miki T., Mizobuchi K.,
RA Mori H., Motomura K., Nakamura Y., Nishimoto H., Nishio Y., Saito N.,
RA Samesi G., Seki Y., Tagami H., Takemoto K., Wada C., Yamamoto Y.,
RA Yano M., Horiiuchi T.;
RT "A 718-kb DNA sequence of the Escherichia coli K-12 genome
RT corresponding to the 12.7-28.0 min region on the linkage map."
RL DNA Res. 3:137-155(1996).
[4]
RP DOMAINS.
RX MEDLINE=91296736; PubMed=2068069;
RA Levensgood S.K., Beyer W.F. Jr., Webster R.E.;
RT "TOL A: a membrane protein involved in colicin uptake contains an
RT extended helical region."
RL Proc. Natl. Acad. Sci. U.S.A. 88:5939-5943(1991).
[5]
RP INTERACTION WITH PORINS.
RX MEDLINE=97133271; PubMed=8978668;
RA Derouiche R., Gavaioli M., Benedetti H., Prilipov A., Lazdunski C.,
RA Llobes R.;
RT "TOL A central domain interacts with Escherichia coli porins."
RL EMBO J. 15:6408-6415(1996).
[6]
RP X-RAY CRYSTALLOGRAPHY (1.85 ANGSTROMS) OF 238-421.
RX MEDLINE=99332679; PubMed=10404600;
RA Lubkowski J., Hennecke F., Plueckhuhn A., Wlodawer A.;
RT "Filamentous phage infection: crystal structure of g3p in complex
RT with its coreceptor, the C-terminal domain of TolA."
RL Structure 7:711-722(1999).
-1- FUNCTION: INVOLVED IN THE TONB-INDEPENDENT UPTAKE OF GROUP A
COLICINS (COLICINS A, EI, E2, E3, AND K). NECESSARY FOR THE
BINDING TO THE BACTERIA. ALSO INVOLVED IN THE TRANSLOCATION
OF BACTERIOPHAGE DNA.
-1- SUBUNIT: INTERACTS, VIA DOMAIN II, WITH PORINS OMPC, OMPF, PHOE
AND LAMB.

CC -1- SUBCELLULAR LOCATION: Type II membrane protein. Inner membrane.
CC -----
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CC -----
DR EMBL: M28232; AAA24683.1; -
DR EMBL: AE000177; AAC73833.1; -
DR EMBL: D90713; BAA35405.1; -
DR PIR: JVO057; JVO057.
DR PDB: 1TOL; 20-MAY-99.
DR Ecogene: BG11007; tolA.
KW Transport; Protein transport; Bacteriocin transport; Transmembrane;
KW Repeat; Inner membrane; 3D-structure; Complete proteome.
FT DOMAIN 1 13 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 14 34 POTENTIAL.
FT DOMAIN 35 421 PERIPLASMIC (POTENTIAL).
FT DOMAIN 48 310 DOMAIN II (ALPHA-HELICAL).
FT DOMAIN 311 421 DOMAIN III (FUNCTIONAL).
FT DOMAIN 224 278 10 X TANDEN REPEATS OF [ED]-K(1,2)-
A(2,4).
FT HELIX 335 349
FT TURN 350 351
FT TURN 353 354
FT HELIX 355 358
FT TURN 359 360
FT STRAND 363 369
FT TURN 371 372
FT STRAND 375 383
FT HELIX 385 397
FT HELIX 406 412
FT TURN 413 414
FT STRAND 416 421
SQ SEQUENCE 421 AA; 43156 MW; 8B2F52BA97C655E CRC64;
Query Match 46.0%; Score 98; DB 1; Length 421;
Best Local Similarity 65.3%; Pred. No. 0.0073;
Matches 32; Conservative 2; Mismatches 9; Indels 6; Gaps 3;
QY 2 KKYAKKAAEQA--KK--AYKAAKQAATYERAAKAAKAAEAYAA 45
DB 247 KKAARAKAAERAAADKKAARAKAAADKRAA-AKAAARAKAAAKAAEAA 294
RESULT 3
ID M2T2_DROHY STANDARD; PRT; 1391 AA.
AC Q08696;
DT 01-FEB-1995 (Rel. 31, Created)
DT 01-FEB-1995 (Rel. 31, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Axoneme-associated protein mst101(2).
GN MST101(2).
OS Drosophila hydei (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydriidae; Drosophilidae; Drosophila.
OX NCBI_TaxId=7224;
[1]
RP SEQUENCE FROM N.A., AND CHARACTERIZATION.
RX MEDLINE=95045538; PubMed=7957199;
RA Neesen J., Padmanabhan S., Buenemann H.;
RT "Tandemly arranged repeats of a novel highly charged 16-amino-acid
RT motif representing the major component of the sperm-tail-specific
RT axoneme-associated protein family Dmst101 form extended
RT alpha-helical rods within the extremely elongated spermatozoa of
RT Drosophila hydei."
RL Eur. J. Biochem. 225:1089-1095(1994).
-1- FUNCTION: POSSIBLE STRUCTURAL ROLE IN THE SPERM TAIL.


```

DR SMART; SMO0411; BHL; 1.
DR POSITE; PS00045; HISTONE LIKE; 1.
KW DNA-binding; DNA condensation; Repeat.
FT DOMAIN 1 90 BACTERIAL HISTONE-LIKE DOMAIN.
FT DOMAIN .101 205 DEGENERATE REPEATS REGION.
SQ SEQUENCE 208 AA; 21230 MW; CAsF577F61F7EF09 CRC64;

Query March 42.0%; Score 89.5; DB 1; Length 208;
Best Local Similarity 53.7%; Pred. No. 0.025;
Matches 29; Conservative 3; Mismatches 13; Indels 9; Gaps 2

OY 1 AKKYAKTAKAKA-----KKAYKAEAKKAKY--EKAARAKAAKEAAYEA 45
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 111 AKKAAKAPAKKAAAKKTATGAAKKAPAKKAAATKAPAKKATATAPAKKAAATKA 164

RESULT 5
H1B STRPU STANDARD; PRT; 211 AA.
AC P1569;
DT 01-APR-1990 (Rel. 14, Created)
DT 01-APR-1990 (Rel. 14, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE Histone H1-beta, late embryonic.
OS Strongylocentrotus purpuratus (Purple sea urchin).
OC Eukaryota; Metazoa; Echinodermata; Eleutherozoa; Echinozoa;
OC Echinoidae; Euechinoidae; Echinacea; Strongylocentrotidae;
OC Strongylocentrotus.
OX NCBI_TaxID=7668;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=68246461; PubMed=2837660;
RA Lai Z.-C., Childs G.;
RT "Characterization of the structure and transcriptional patterns of
RT the gene encoding the late histone subtype H1-beta of the sea urchin
RT Strongylocentrotus purpuratus."
RL Mol. Cell. Biol. 8:1842-1844(1988).

-1- FUNCTION: HISTONES H1 ARE NECESSARY FOR THE CONDENSATION OF
CC NUCLEOSOME CHAINS INTO HIGHER ORDER STRUCTURES.
CC -1- SUBCELLULAR LOCATION: Nuclear.
CC -1- SIMILARITY: BELONGS TO THE HISTONE H1/H5 FAMILY.
CC -----
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CC -----
CC EMBL; M20314; AAA30052.1; -.
DR PIR; A28100; A28100.
DR HSBP; P02259; IHST.
DR InterPro; IPR005818; Histone_H1/H5.
DR InterPro; IPR005819; Histone_H5.
DR InterPro; IPR003216; Linkerhist_N.
DR Pfam; PF00538; linker_histone; 1.
DR PRINTS; PR00624; HISTONH5.
DR ProDom; PD000373; linkerhist_N; 1.
DR SMART; SMO0526; H15. 1.
DR K0 Chromosomal protein; Nuclear protein; DNA-binding; Multigene family.
SQ SEQUENCE 211 AA; 22169 MW; 9F214581334BBE7A CRC64;

Query March 42.0%; Score 89.5; DB 1; Length 211;
Best Local Similarity 61.9%; Pred. No. 0.026;
Matches 26; Conservative 3; Mismatches 10; Indels 3; Gaps 2

OY 1 AKKYAKTAKAKAKKAYKAAEAKKAAATYKAAAEKAAAEKAA 42
      :||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 139 SKTTTKVKVKKPAKKAKKPA--AKKAAK--KPAKKKPAKAA 177

RESULT 6

```

ID	DBH_MVCTU	STANDARD:	PRT:	214 AA.
AC	P95109;			
DT	15-JUL-1999	(Rel. 38, Created)		
DT	15-JUL-1999	(Rel. 38, Last sequence update)		
DT	16-OCT-2001	(Rel. 40, Last annotation update)		
DE	DNA-binding protein HU homolog (Histone-like protein) (Htp) (21-kDa			
DE	lamini-2-binding protein).			
DN	HUP OR HLP OR LBP21 OR RV2986C OR MT3064 OR MTCY349.01.			
OS	Mycobacterium tuberculosis.			
OC	Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;			
OC	Corynebacteriaceae; Mycobacteriaceae; Mycobacterium.			
OX	NCBI_TaxID=1773;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	STRAIN=H37Rv;			
RC	MEDLINE=968295987; PubMed=9634230;			
RA	Cole S.T., Broch R., Parkhill J., Garnier T., Churcher C., Harris D.,			
RA	Gordon S.V., Eiglmeier K., Gae S., Barry C.E. III, Tekela F.,			
RA	Badcock K., Bisham D., Brown D., Chillingworth T., Connor R.,			
RA	Davies R., Devlin K., Fellwell T., Gentles S., Hamlin N., Holroyd S.,			
RA	Holmesby T., Jorgels K., Krogh A., McLean J., Moule S., Murphy L.,			
RA	Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,			
RA	Rutter S., Seeger K., Skelton S., Squares S., Squares R.,			
RA	Sulston J.B., Taylor K., Whitehead S., Barrett B.G.,			
RT	"Deciphering the biology of Mycobacterium tuberculosis from the			
RT	complete genome sequence.";			
RL	Nature 393:537-544 (1998).			
RN	[2]			
RP	SEQUENCE FROM N.A.			
RC	STRAIN=CDC 1551 / Oshkosh;			
RA	Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,			
RA	Petersen J., Deboy R., Dodson R., Gwinn M.L., Haft D., Hickey E.,			
RA	Kolony J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L.,			
RA	Delcher A., Utterback T., Weidman J., Khouri H., Gill J., Mikula A.,			
RA	Bisler W.;			
RT	"Whole genome comparison of Mycobacterium tuberculosis clinical and			
RT	laboratory strains.";			
RL	Submitted (Apr-2001) to the EMBL/GenBank/DBJ databases.			
RN	[3]			
RP	SEQUENCE OF 71-86, AND DNA-BINDING.			
RC	STRAIN=H37Rv;			
RA	Prasad H.K., Annapuram P.S., Dey A.B., Tyagi J.S., Jain N.K.,			
RA	Saxena P.;			
RL	Submitted (DEC-1997) to the SWISS-PROT data bank.			
CC	-1- FUNCTION: THIS PROTEIN BELONGS TO THE HISTONE LIKE FAMILY OF			
CC	PROKARYOTIC DNA-BINDING PROTEINS WHICH ARE CAPABLE OF WRAPPING			
CC	DNA TO STABILIZE IT, AND PREVENT ITS DENATURATION UNDER EXTREME			
CC	ENVIRONMENTAL CONDITIONS (BY SIMILARITY).			
CC	-1- SIMILARITY: BELONGS TO THE BACTERIAL HISTONE-LIKE PROTEIN FAMILY.			
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CC	or send an email to license@sib-sib.ch).			
CC	-----			
CC	EMBL; Z83018; CAB05427.1; -			
DR	EMBL; AE007127; AAK47393.1; -			
DR	PIR; G70673; G70673.			
DR	HSBP; P02346; 1HTU.			
DR	TIGR; MT3064; -			
DR	TubercuLib; RV2986C; -			
DR	InterPro; IPR000119; Bac DNABind.			
DR	InterPro; IPR005819; Histone H5.			
DR	Pfam; PF00216; Bac DNA binding; 1.			
DR	PRINTS; PR00624; HISTONEH5.			
DR	Prodom; PD000945; Bac DNABind; 1.			
DR	SMART; SM00411; BHL_1.			
DR	PROSITE; PS00045; HISTONE_LIKE; 1.			
KW	DNA-binding; DNA condensation; Repat; Complete proteome.			

FT	DOMAIN	1	90	BACTERIAL HISTONE-LIKE DOMAIN.
FT	DOMAIN	101	214	DEGENERATE REPEATS REGION.
SEQ	SEQUENCE	214 AA;	22187 MM;	CB09AF20FB353544 CRC64;
Query Match	Similarity	42.0%;	Score 89.5;	DB 1; Length 214;
Best Local	Similarity	55.6%;	Pred. No. 0.026;	
Matches	25;	Conservative	4;	Mismatches 13; Indels 3; Gaps 1
DB	111 AKYAKAKAPAKKATKAKAKATKAP	---	RCATKAPAKKATATYA	152
QY	1 AKYAKAKAKAKAKAKAYKAAEAKKAKYKKAAYKAAEAKKAAEAYEA	45		
RESULT 7				
ID	TMPB	TREPB	STANDARD;	PRT; 384 AA.
AC	P29720;			
DT	01-APR-1993	(Rel. 25, Created)		
DT	01-APR-1993	(Rel. 25, Last sequence update)		
DT	16-OCT-2001	(Rel. 40, Last annotation update)		
DB	Treponema	membrane protein B precursor (Antigen tmpb).		
OS	TMPB.			
GN	Treponema phagedentis.			
OC	Bacteria; Spirochaetes; Spirochaetales; Spirochaetaceae; Treponema.			
OX	NCBI_Textid=162;			
KN	(1)			
RN	SEQUENCE FROM N.A.			
RC	STRAIN=Kazan 5;			
RX	MEDLINE=91372983; PubMed=1894368;			
RA	Yelton D.B., Limberger R.J., Curci K., Malinovsky-Rumel F.,			
RA	Slivensky L., Schouls L.M., van Embden J.D., Charon N.W.;			
RT	"Treponema phagedentis encodes and expresses homologs of the Treponema			
RL	pallidum tmpa and tmpb proteins."			
CC	Infect. Immun. 59:3685-3693 (1991).			
CC	-1- FUNCTION: TMP MAY SERVE AS A PORIN OR TRANSPORT PROTEIN FOR			
CC	LARGE MOLECULES.			
CC	-1- SUBCELLULAR LOCATION: OUTER MEMBRANE-ASSOCIATED.			
CC	-1- SIMILARITY: TO TMPB OF T.PALLIDUM.			
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CC	or send an email to license@isb-sib.ch).			
CC	-----			
DR	EMBL; M58563; AAA27480.1; -.			
DR	PIR; B43592; B43592.			
KW	Antigen; Outer membrane; Repeat; Signal.			
FT	SIGNAL	1	21	POTENTIAL.
FT	CHAIN	22	384	TREPONEMA MEMBRANE PROTEIN B.
FT	DOMAIN	151	235	17 X 5 AA TANDEM REPEATS OF K-A-A-(AKR)-
FT				(ED).
FT	REPEAT	151	155	1-1.
FT	REPEAT	156	160	1-2.
FT	REPEAT	161	165	1-3.
FT	REPEAT	166	170	1-4.
FT	REPEAT	171	175	1-5.
FT	REPEAT	176	180	1-6.
FT	REPEAT	181	185	1-7.
FT	REPEAT	186	190	1-8.
FT	REPEAT	191	195	1-9.
FT	REPEAT	196	200	1-10.
FT	REPEAT	201	205	1-11.
FT	REPEAT	206	210	1-12.
FT	REPEAT	211	215	1-13.
FT	REPEAT	216	220	1-14.
FT	REPEAT	221	225	1-15.
FT	REPEAT	226	230	1-16.
FT	REPEAT	231	235	1-17.
FT	DOMAIN	236	288	6 X 6 AA TANDEM REPEATS OF [EA]-A-A-R-X-
FT				A-A-E.


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CC -1- PTM: GLYCOSYLATED. GLYCOSYLATION MAY PROTECT THE PROTEIN FROM
CC PROTEOLYTIC DEGRADATION AND BE IMPORTANT FOR HEMAGGLUTINATION. IT
CC SUGGESTS THAT THE CARBOHYDRATE MOIETY MAY BE LOCATED WITHIN THE C-
CC TERMINAL DOMAIN OF HBHA.
CC -1- MISCELLANEOUS: SERUM FROM PATIENTS DIAGNOSED WITH ACTIVE
CC TUBERCULOSIS THAT HAD NOT BEEN VACCINATED CONTAINS ANTIBODIES THAT
CC RECOGNIZE HBHA, WHEREAS SERUM FROM HEALTHY INDIVIDUALS DOES NOT
CC CONTAIN ANY.
CC -1- SIMILARITY: STRONG, TO M. LEPRAE HBHA.
CC -----
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CC -----
CC EMBL, AF074390; AAC26052.1; -.
CC DR EMBL; 277162; CAB00936.1; -.
CC DR EMBL; AE006951; AAK44716.1; ALT INIT.
CC DR PIR; F70742; F70742.
CC TIGR; MT0493; -.
CC DR Tuberculin; K0475; -.
CC DR Cell adhesion; Heparin-binding; Hemagglutinin; Glycoprotein;
CC KM Virulence; Complete proteome.
CC FT INIT MET 0
CC FT DOMAIN 151 193 ALA/LYS-RICH.
CC FT CONFLICT 120 120 R -> P (IN REF. 1).
CC FT SEQUENCE 198 AA, 21403 MW, 51376066F1EB6042 CRC64;
CC SQ
CC
CC Query Match 40.4%; Score 86; DB 1; Length 198;
CC Best Local Similarity 65.6%; Pred. No. 0.053;
CC Matches 21; Conservative 4; Mismatches 7; Indels 0; Gaps 0;
CC
CC QY 11 EKAKKAYRAAEKKAAYEKAARAKAEEA 42
CC Db 160 KKAAPAKKAAPAKKAAPAKKAAPAKKA 191
CC
CC RESULT 10
CC MST1_DROXY STANDARD; PRT; 344 AA.
CC AC Q08695;
CC DT 01-FEB-1995 (Rel. 31, Created)
CC DT 01-FEB-1995 (Rel. 31, Last sequence update)
CC DT 16-OCT-2001 (Rel. 40, Last annotation update)
CC DE Axoneme-associated protein mst101(1).
CC GN MST101(1).
CC OS Drosophila hydei (Fruit fly).
CC OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
CC OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
CC OC Ephydroidea; Drosophilidae; Drosophila.
CC OC NCBI_Taxid=7224;
CC RN [1]
CC RP SEQUENCE FROM N.A., AND CHARACTERIZATION.
CC RC TISSUE=Testis;
CC RX MEDLINE=94200512; PubMed=8150205;
CC RA Neesen J., Buenen H., Heinlein U.A.;
CC RT "The Drosophila hydei gene Dmst101(1) encodes a testis-specific,
CC RT repetitive, axoneme-associated protein with differential abundance in
CC RT Y chromosomal deletion mutant flies.";
CC RL Dev. Biol. 162:414-425 (1994).
CC CC -1- FUNCTION: POSSIBLE STRUCTURAL ROLE IN THE SPERM TAIL. IT IS
CC CC ASSOCIATED WITH AXONEMAL STRUCTURES.
CC CC -1- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC CC -1- TISSUE SPECIFICITY: TESTIS. LOCATED IN SPERMATOCYTES AND
CC CC SPERMATID BUNDLES.
CC CC -1- DOMAIN: THE PREDOMINANT STRUCTURE IS ALPHA-HELICAL.
CC CC -1- POLYMORPHISM: THE NUMBER OF REPEATS VARIES BETWEEN STRAINS.
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CC -----
CC EMBL, X73480; CA51875.1; -.
CC DR PIR; S34153; S34153.
CC DR FlyBase; FBgn0011816; Dhyd\mst101(1).
CC KW Sperm; Repeat; Multigene family.
CC FT DOMAIN 58 337
CC FT SEQUENCE 344 AA, 37793 MW, 24C65D2510387E2A CRC64;
CC SQ
CC
CC Query Match 40.4%; Score 86; DB 1; Length 344;
CC Best Local Similarity 57.8%; Pred. No. 0.087;
CC Matches 26; Conservative 3; Mismatches 12; Indels 4; Gaps 2;
CC
CC QY 2 KKYAKKAR-AEKAKKAYKAERK---AAKYEKAARAKAEEA 42
CC Db 69 KEAAEKKCAAPAKKAEKKAERKCAEAKEKKAERKCAEA 113
CC
CC RESULT 11
CC TOLA_PSEAE STANDARD; PRT; 347 AA.
CC AC P50600;
CC DT 01-OCT-1996 (Rel. 34, Created)
CC DT 16-OCT-2001 (Rel. 40, Last sequence update)
CC DT 16-OCT-2001 (Rel. 40, Last annotation update)
CC DE Tola protein.
CC GN TOLA OR PA0971.
CC OS Pseudomonas aeruginosa.
CC OC Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;
CC OC Pseudomonadaceae; Pseudomonas.
CC OC NCBI_Taxid=287;
CC RN [1]
CC RP SEQUENCE FROM N.A.
CC RC STRAIN=PAO;
CC RX MEDLINE=97113525; PubMed=8955385;
CC RA Dennis J.J., Lafontaine E.R., Sokol P.A.;
CC RT "Identification and characterization of the tolR genes of
CC RT Pseudomonas aeruginosa.";
CC RL J. Bacteriol. 178:7059-7068 (1996).
CC RN [2]
CC RP REVISIONS TO N-TERMINUS.
CC RA Duan K., Sokol P.A.;
CC RL Submitted (AUG-1999) to the EMBL/GenBank/DBJ databases.
CC RN [3]
CC RP SEQUENCE FROM N.A.
CC RC STRAIN=ATCC 15692 / PAO1;
CC RX MEDLINE=20437337; PubMed=10984043;
CC RA Stover C.K., Pham X.-Q.T., Erwin A.L., Mizoguchi S.D., Warren P.,
CC RA Hickey M.J., Brinkman F.S.L., Hufnagle W.O., Kowalik D.J., Lagrou M.,
CC RA Garber R.L., Goltzy L., Tolentino E., Westbrook-Wadman S., Yuan Y.,
CC RA Brody L.L., Coulter S.N., Folger K.R., Kas A., Labdig K., Lim R.M.,
CC RA Smith K.A., Spencer D.H., Wong G.K.-S., Wu Z., Paulsen I.T.,
CC RA Reizer J., Sater M.H., Hancock R.E.W., Lory S., Olson M.V.;
CC RT "Complete genome sequence of Pseudomonas aeruginosa PAO1, an
CC RT opportunistic pathogen.";
CC RL Nature 406:959-964 (2000).
CC CC -1- FUNCTION: INVOLVED IN THE TONB-INDEPENDENT UPTAKE OF PROTEINS
CC CC (BY SIMILARITY).
CC CC -1- SUBCELLULAR LOCATION: Type II membrane protein. Inner membrane
CC CC (potential).
CC CC -----
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CC -----
DR EMBL; U39558; AAC4660.2; -.
DR EMBL; AB004530; AAG04360.1; -.
DR PIR; E83525; E83525.
DR InterPro; IPR006260; ToMB_C.
DR TIGRfam; TIGR01352; ToMB_Cterm; 1.
KW Transport; Protein transport; Transmembrane; Repeat; Inner membrane;
KM Complete proteome.
FT DOMAIN 1 16 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 17 37 POTENTIAL.
FT DOMAIN 38 347 PERIPLASMIC (POTENTIAL).
FT DOMAIN 209 216 POLY-ALA.
SQ SEQUENCE 347 AA; 37935 MW; EEDD4B04AA095945 CRC64;

Query Match 40.4%; Score 86; DB 1; Length 347;
Best Local Similarity 51.1%; Pred. No. 0.088;
Matches 24; Conservative 6; Mismatches 15; Indels 2; Gaps 1;

OY 1 AKKYAKKAKKAKKAKK--YKAAEAKKAKYKAAEAKKAAEAKAAE 45
DB 125 AOKAAEAKKADKAAEAKKAEKQKQADIAKGRABEDAKKAAEADA 171

RESULT 12
DBH MYCBO STANDARD; PRT; 205 AA.
AC Q9XB18; Q9S5J5;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE DNA-binding protein HU homolog (Histone-like protein) (HLP).
GN HUP OR HLP OR MDP1.
OS Mycobacterium bovis.
OC Bacteria; Actinobacteria; Actinomycetales;
OC Corynebacteriaceae; Mycobacteriaceae; Mycobacterium.
OX NCBI_TaxID=1765;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ANS;
RA Prabhakar S., Tyagi J.S., Praead H.K.;
RT "HMPM-A target for differentiation of M.tuberculosis and M.bovis.";
RL Submitted (NOV-1998) to the EMBL/Genbank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=BGC / Tokyo;
RA Matsumoto S., Yukioka H., Matsuo T., Mineda T., Yamada T.;
RT "Identification of a novel protein generating bacterial slow growth
from Mycobacterium.";
RL Submitted (APR-1998) to the EMBL/Genbank/DBJ databases.
CC -1- FUNCTION: THIS PROTEIN BELONGS TO THE HISTONE LIKE FAMILY OF
PROKARYOTIC DNA-BINDING PROTEINS WHICH ARE CAPABLE OF WRAPPING
DNA TO STABILIZE IT, AND PREVENT ITS DENATURATION UNDER EXTREME
ENVIRONMENTAL CONDITIONS (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE BACTERIAL HISTONE-LIKE PROTEIN FAMILY.
CC -----
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CC -----
DR EMBL; Y18421; CAB46493.1; -.
DR EMBL; AB013441; BAA78330.1; -.
DR HSSP; P02346; 1HUU.
DR InterPro; IPR000119; Bac_DNAbind.
DR InterPro; IPR005819; Histone_H5.
DR Pfam; PF00216; Bac_DNA_binding; 1.
DR PRINTS; PR00624; HISTONH5.
DR ProDom; PD000945; Bac_DNAbind; 1.
DR SMART; SM00411; BHL; 1.
DR PROSITE; PS00045; HISTONE_LIKE; 1.

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KW DNA-binding; DNA condensation; Repeat.
FT DOMAIN 1 90 BACTERIAL HISTONE-LIKE DOMAIN.
FT DOMAIN 101 205 DEGENERATE REPEATS REGION.
FT CONFLICT 199 199 A -> T (IN REF. 2).
SQ SEQUENCE 205 AA; 21262 MW; 19FC667885DPE6A8 CRC64;

Query Match 40.1%; Score 85.5; DB 1; Length 205;
Best Local Similarity 58.5%; Pred. No. 0.061;
Matches 24; Conservative 2; Mismatches 12; Indels 3; Gaps 1;

OY 1 AKKYAKKAKKAKKAKKAYKAAEAKKAKYKAAEAKKAAEAKA 41
DB 111 AKKYAKKAPAKKATKAAKKAATKAPAA--KKAATKAPAKKA 148

RESULT 13
RS16 CORGL STANDARD; PRT; 165 AA.
AC Q8NNX3;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE 30S ribosomal protein S16.
GN RSP OR CGL2054.
OS Corynebacterium glutamicum (Brevibacterium flavum).
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Corynebacteriaceae; Corynebacteriaceae; Corynebacterium.
OX NCBI_TaxID=1718;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 13032 / DSM 20300 / NCIB 10025;
RA Nakagawa S.;
RT "Complete genomic sequence of Corynebacterium glutamicum ATCC 13032.";
RL Submitted (MAY-2002) to the EMBL/Genbank/DBJ databases.
CC -1- SIMILARITY: BELONGS TO THE S16P FAMILY OF RIBOSOMAL PROTEINS.
CC -----
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CC -----
DR EMBL; AP005280; BAB99447.1; -.
DR HAMAP; MF_00385; -; 1.
DR InterPro; IPR000307; Ribosomal_S16.
DR Pfam; PF00886; Ribosomal_S16; 1.
DR ProDom; PD003791; Ribosomal_S16; 1.
DR TIGRfam; TIGR00002; S16; 1.
DR PROSITE; PS00732; RIBOSOMAL_S16; FALSE_NEG.
KW Ribosomal protein; Complete proteome.
SQ SEQUENCE 165 AA; 17837 MW; 61DD81961BC30846 CRC64;

Query Match 38.5%; Score 82; DB 1; Length 165;
Best Local Similarity 62.9%; Pred. No. 0.11;
Matches 22; Conservative 4; Mismatches 7; Indels 2; Gaps 1;

OY 7 KAKAEKAKKAYKAAEAKKAKYKAAEAKKAAEAKA 41
DB 120 EAITEKKKKAEDEKAEKAA--EKAAEAKAADAAS 152

RESULT 14
H1 PARAN STANDARD; PRT; 248 AA.
AC P03256;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE Histone H1, gonadal.
OS Parachinus angulosus (Angulate sea urchin).
OC Eukaryota; Metazoa; Echinodermata; Eleutherozoa; Echinozoa;

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CC Echinoidea; Echinoidea; Echinacea; Echinoidea; Echinidae;
CC Parechinus.
OX NCBI TaxId=7658;
RN [1]
RP SEQUENCE OF 1-84.
RX MEDLINE=80156831; PubMed=6767609;
RA Strickland W.N., Strickland M., de Groot P.C., von Holt C.,
RA Wiltmann-Liebold B.;
RT "The primary structure of histone H1 from sperm of the sea urchin
RT Parechinus angulosus. 1. Chemical and enzymatic fragmentation of the
RT protein and the sequence of amino acids in the four N-terminal
RT cyanogen bromide peptides."
RL Eur. J. Biochem. 104:559-566 (1980).
RN [2]
RP SEQUENCE OF 80-248.
RX MEDLINE=80156832; PubMed=7363905;
RA Strickland W.N., Strickland M., Brandt W.F., von Holt C., Lehmann A.,
RA Wiltmann-Liebold B.;
RT "The primary structure of histone H1 from sperm of the sea urchin
RT Parechinus angulosus. 2. Sequence of the C-terminal CNBr peptide and
RT the entire primary structure."
RL Eur. J. Biochem. 104:567-578 (1980).
CC -1- FUNCTION: HISTONES H1 ARE NECESSARY FOR THE CONDENSATION OF
CC NUCLEOSOME CHAINS INTO HIGHER ORDER STRUCTURES.
CC -1- SUBCELLULAR LOCATION: Nuclear.
CC -1- TISSUE SPECIFICITY: Sperm.
CC -1- SIMILARITY: BELONGS TO THE HISTONE H1/H5 FAMILY.
DR HSSP; P02259; HST.
DR InterPro; IPR005818; Histone_H1/H5.
DR InterPro; IPR005819; Histone_H5.
DR InterPro; IPR003216; Linkerhist_N.
DR Pfam; PF00538; Linker histone; 1.
DR PRINTS; PR00624; HISTONEH5.
DR ProDom; PD000373; Linkerhist_N; 1.
DR SMART; SM00526; H15; 1.
DR Chromosomal protein; Nuclear protein; DNA-binding; Multigene family;
KW Sperm.
KM
KV
FT VARIANT 144 144 K -> R.
SQ SEQUENCE 248 AA; 26387 MW; 1B25B3F136541947 CRC64;

Query Match 38.5%; Score 82; DB 1; Length 248;
Best Local Similarity 57.8%; Pred. No. 0.16;
Matches 26; Conservative 3; Mismatches 14; Indels 2; Gaps 2;

OY 2 KKYAKKAKAKAKAKYKAAEAKKAKYKAAEAKKAA-AKAAAYEA 45
DB 120 KKAATTSAAAKAKA-KAAAKKAKAKAKAAKKAALAKKAAA 163

RESULT 15
HIE CHIPA STANDARD; PRT; 235 AA.
AC P40762;
DT 01-FEB-1995 (rel. 31, Created)
DT 01-FEB-1995 (rel. 31, Last sequence update)
DT 15-JUL-1999 (rel. 38, Last annotation update)
DE Histone H1B.
OS Chironomus pallidivittatus (Midge).
OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
OC Neoptera; Endopterygota; Diptera; Nematocera; Chironomidae;
OC Chironomidae; Chironominae; Chironomus.
OX NCBI TaxId=7151;
RN [1]
RP SEQUENCE FROM N.A.
RA Schulze E., Mieniewicz J.R., Nagel S., Gavenis K., Grosbach U.;
RL Submitted (XXL-1994) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: HISTONES H1 ARE NECESSARY FOR THE CONDENSATION OF
CC NUCLEOSOME CHAINS INTO HIGHER ORDER STRUCTURES.
CC -1- SUBCELLULAR LOCATION: Nuclear.
CC -1- SIMILARITY: BELONGS TO THE HISTONE H1/H5 FAMILY.
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CC -----
DR EMBL; U29106; AAA21714.1; -.
DR HSSP; P08287; IGHC.
DR InterPro; IPR005818; Histone_H1/H5.
DR InterPro; IPR005819; Histone_H5.
DR InterPro; IPR003216; Linkerhist_N.
DR Pfam; PF00538; Linker histone; 1.
DR PRINTS; PR00624; HISTONEH5.
DR ProDom; PD000373; Linkerhist_N; 1.
DR SMART; SM00526; H15; 1.
DR Chromosomal protein; Nuclear protein; DNA-binding; Multigene family.
KV
SQ SEQUENCE 235 AA; 24446 MW; 340BC5B9A85002AC CRC64;

Query Match 38.0%; Score 81; DB 1; Length 235;
Best Local Similarity 41.3%; Pred. No. 0.19;
Matches 26; Conservative 2; Mismatches 17; Indels 18; Gaps 1;

OY 1 AKKYAKKAKAKA-----KKAAYKAAEAKKAKYKAAEAKKAAKAA 42
DB 126 AKKVDKPKAPAPAPKPKSTNRVTGKKVTKKPAKKPKKATYAKAPKAAKAPAA 185

OY 43 YEA 45
DB 186 KKA 188

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Search completed: January 21, 2004, 09:00:58
 Job time : 8.13636 secs

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OM protein - protein search, using sw model

Run on: January 21, 2004, 08:53:31 ; Search time 18.1169 Seconds
(without alignments)
640,969 Million cell updates/sec

Title: US-09-816-989A-2

Perfect score: 213
Sequence: 1 AKKAKKAKAKAKAKAYKAA.....AKYKAAAKKAAKAAAYEA 45

Scoring table:
BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 830525 seqs, 258052604 residues

Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

SPTREMBL 23:*

- 1: sp_archaea:*
- 2: sp_bacteria:*
- 3: sp_fungi:*
- 4: sp_human:*
- 5: sp_invertebrate:*
- 6: sp_mammal:*
- 7: sp_nhc:*
- 8: sp_organelle:*
- 9: sp_plant:*
- 10: sp_rodent:*
- 11: sp_virus:*
- 12: sp_vertebrate:*
- 13: sp_unclassified:*
- 14: sp_virus:*
- 15: sp_bacteriopl:*
- 16: sp_bacteriopl:*
- 17: sp_archaea:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	98	46.0	394	16	Q8X965
2	98	46.0	421	16	Q8FJ11
3	94.5	44.4	239	16	Q8V5M4
4	91	42.7	244	16	Q8AJX2
5	90	42.3	243	16	Q8ZAE7
6	88.5	41.5	275	5	Q01395
7	88.5	41.5	372	2	Q9WXX1
8	88	40.3	899	16	Q9ASJ6
9	86.5	40.6	200	16	Q8XVW7
10	86	40.4	155	16	Q8PI40
11	86	40.4	168	16	Q8F930
12	86	40.4	212	3	Q93946
13	85	39.9	98	5	Q8WQ44
14	85	39.9	111	5	Q8TR93
15	85	39.9	182	2	Q45370
16	84	39.4	60	5	Q9U3W3

17	84	39.4	144	13	Q8AW13	Q8AW13 oncorhynch
18	83.5	39.2	483	12	Q8ONDI	Q8ONDI octocarpus
19	83	39.0	293	10	Q9AT18	Q9AT18 lens culina
20	83	39.0	389	16	Q9CM70	Q9CM70 pasteurella
21	82.5	38.7	243	5	Q23784	Q23784 chironomus
22	82	38.5	190	5	Q8MYC3	Q8MYC3 mytilus cal
23	82	38.5	262	16	Q9S2M2	Q9S2M2 streptomyce
24	82	38.5	568	3	Q94567	Q94567 schizosacch
25	81.5	38.3	156	16	Q8PEU0	Q8PEU0 xanthomonas
26	81.5	38.3	311	12	Q84528	Q84528 paramacium
27	81.5	38.3	482	2	Q93LK4	Q93LK4 enterococcu
28	81	38.0	81	5	Q9NFP6	Q9NFP6 trypanosoma
29	81	38.0	112	5	Q9XYR5	Q9XYR5 leishmania
30	81	38.0	241	5	Q23790	Q23790 chironomus
31	81	38.0	290	10	Q9AT24	Q9AT24 pismu saativ
32	81	38.0	295	10	Q8LKH9	Q8LKH9 pismu fulvu
33	81	38.0	295	10	Q8LKH0	Q8LKH0 pismu saativ
34	81	38.0	295	10	Q9AT22	Q9AT22 lathyrus sa
35	81	38.0	295	10	Q9ZK20	Q9ZK20 pismu saativ
36	81	38.0	295	10	Q9AT25	Q9AT25 pismu saativ
37	81	38.0	297	10	Q9SXQ8	Q9SXQ8 pismu saativ
38	81	38.0	298	10	Q8LKH1	Q8LKH1 lathyrus ap
39	81	38.0	301	10	Q9AT23	Q9AT23 pismu saativ
40	81	38.0	306	10	Q9AT21	Q9AT21 lathyrus sa
41	80.5	37.8	277	16	Q9XAO3	Q9XAO3 streptomyce
42	80.5	37.8	445	3	Q9P3Q8	Q9P3Q8 neurospora
43	80	37.6	66	5	Q95QZ0	Q95QZ0 caenorhabdi
44	80	37.6	71	5	Q9NF08	Q9NF08 trypanosoma
45	80	37.6	81	5	Q9N6L9	Q9N6L9 trypanosoma

ALIGNMENTS

RESULT 1

ID Q8X965 PRELIMINARY; PRT; 394 AA.

AC Q8X965.

DT 01-MAR-2002 (TREMBLrel. 20, Last sequence update)

DT 01-OCT-2002 (TREMBLrel. 22, Last annotation update)

DE Membrane spanning protein, required for outer membrane integrity

DE (Membrane spanning protein Tola).

GN TOLA OR Z0907 OR EGS0774.

OS Escherichia coli O157:H7.

OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;

OC Enterobacteriaceae; Escherichia.

OX NCI_TaxID=83334;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=O157:H7 / EDI933 / ATCC 700927;

RX MEDLINE=21074935; PubMed=11206551;

RA Rose D.J., Mayhew G.F., Evans P.S., Gregor J., Kirkpatrick H.A.,

RA Postel G., Hackett J., Klink S., Boutin A., Shao Y., Miller L.,

RA Grobbeck E.J., Davis N.W., Lim A., Dimalanta E.T., Potamoukis K.,

RA Apodaca J., Anantharaman T.S., Lin J., Yen G., Schwartz D.C.,

RA Welch R.A., Blattner F.R.;

RT "Genome sequence of enterohaemorrhagic Escherichia coli O157:H7.";

RL Nature 409:529-533(2001).

RN [2]

RP SEQUENCE FROM N.A.

RC STRAIN=O157:H7 / RIMD 0509952;

RX MEDLINE=21156231; PubMed=11258796;

RA Hayashi T., Makino K., Ohnishi M., Kurokawa K., Ishii K., Yokoyama K.,

RA Han C.-G., Ohtsubo E., Nakayama K., Murata T., Tanaka M., Tobe T.,

RA Iida T., Takami H., Honda T., Sasaki C., Ogasawara N., Yasunaga T.,

RA Kuhara S., Shiba T., Hattori M., Shinagawa H.,

RT "Complete genome sequence of enterohaemorrhagic Escherichia coli

RT O157:H7 and genomic comparison with a laboratory strain K-12.";

RL DNA Res. 8:11-22(2001).

DR EMBL; AF005252; AAC55075.1; -

DR EMBL; AF005253; BAB34197.1; -

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RA Jones L. M., Kaerst U., Kreitt J., Kunin M., Kumb F., Kutepkat G.,
RA Madueno E., Maitounam A., Mala Vicente J., Ng E., Nedjari H.,
RA Nordstedt G., Novella S., de Pablo B., Perez-Diaz J.-C., Purcell R.,
RA Remmel B., Rose M., Schlunet T., Słomow N., Tierrez A.,
RA Vazquez-Poland J.-A., Vose H., Wehland J., Cossart P.,
RA "Comparative genomics of Listeria species.",
RL Science 294:849-852(2001).
DR EMBL; AL591981; CAD00019.1; -.
DR ListList; LMO01941; -.
DR InterPro; IPR002482; LSM.
DR Pfam; PF01476; LSM; 1.
DR SMART; SM00257; LysM; 1.
DR Hypothetical protein; Complete proteome.
SQ SEQUENCE 239 AA; 25836 MW; 72E59D576E0D7832 CRC64;

Query Match 44.4%; Score 94.5; DB 16; Length 239;
Best Local Similarity 56.2%; Pred. No. 0.045;
Matches 27; Conservative 7; Mismatches 9; Indels 5; Gaps 2

Qy 2 KCTAKKAAEKAKK--AYKAAEKK--AKYEGAAEKAAKEAYE 44
Db 124 KAAAEKKAADKKKOEBAVVAANKKQGEAAEKRAADKAAAEKAAAE 171

RESULT 4
Q9AJX2 PRELIMINARY; PRT; 244 AA.
AC Q9AJX2;
DT 01-JUN-2001 (TREMBlrel. 17, Created)
DT 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
DE Putative secreted protein.
GN SC01805 OR SC133.04.
OS Streptomyces coelicolor.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Streptomycinae; Streptomycetaceae; Streptomyces.
OC NCBI_TaxID=1902;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=A3(2) / M145.
RX MEDLINE=21996410; PubMed=1200953;
RA Bentley S.D., Chater K.F., Cerdano-Tarrega A.-M., Challis G.L.,
RA Thomson N.R., James K.D., Harris D.E., Quail M.A., Kleiser H.,
RA Harper D., Bateman A., Brown S., Chandra G., Chen C.W., Collins M.,
RA Cronin A., Frazer A., Goble A., Hidalgo J., Hornsby T., Howarth S.,
RA Huang C.-H., Kleiser I., Larke L., Murphy L., Oliver K., O'Neil S.,
RA Rabinowitsch E., Rajandream M.A., Rutherford K., Ruter S.,
RA Seeger K., Saunders D., Sharp S., Squares R., Squares S., Taylor K.,
RA Warren T., Wietzorrek A., Woodward J., Barrell B.G., Parkhill J.,
RA Hopwood D.A.;
RT "Complete genome sequence of the model actinomycete Streptomyces
RT coelicolor A3(2).",
RL Nature 417:141-147(2002).
DR EMBL; AL939110; CAC28545.1; -.
KW Complete proteome.
SQ SEQUENCE 244 AA; 25524 MW; 61999D62CA23A7B0 CRC64;

Query Match 42.7%; Score 91; DB 16; Length 244;
Best Local Similarity 59.5%; Pred. No. 0.1;
Matches 25; Conservative 4; Mismatches 11; Indels 2; Gaps 1

Qy 5 AKKAAEKAKK--AYKAAEKKAAYEKAAEKAAKEAYE 44
Db 78 SQKVAATKAAKAAKATACKAAVEKKAEEKAAEKAAKAAKE 119

RESULT 5
Q92A67 PRELIMINARY; PRT; 243 AA.
AC Q92A67;
DT 01-DEC-2001 (TREMBlrel. 19, Created)
DT 01-DEC-2001 (TREMBlrel. 19, Last sequence update)
DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)

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DE Hypothetical protein lin2055.
 GN LIN2055.
 OS Listeria innocua.
 CC Bacteria; Firmicutes; Bacillales; Listeriaceae; Listeria.
 CX NCBI_TaxID=1642;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=CLIP 11262 / Serovar 6a;
 RX PubMed=11679669;
 RA Glaeser P., Frangeul L., Buchrieser C., Rusniok C., Amend A.,
 Baquero F., Berche P., Bloeker H., Brandt P., Chakraborty T.,
 Charbit A., Checrouni F., Couve E., de Daruvar A., Deloux P.,
 Domann E., Dominguez-Bernal G., Duchaud E., Durant L., Dussurget O.,
 Entian K.-D., Feibi H., Garcia-del Portillo F., Garrido P.,
 Gautier L., Goebel W., Gomez-Lopez N., Hain T., Hauf J., Jackson D.,
 Jones L.-M., Kaerst U., Klett U., Kunz M., Kuntz F., Kurapat G.,
 Madueno E., Maitournam A., Mata Vicente J., Ng E., Nedjari H.,
 Nordstedt G., Novella S., de Pablo B., Perez-Diaz J.-C., Purcell R.,
 Remmel B., Rose M., Schluter T., Simoes N., Tierrez A.,
 Vazquez-Boland J.-A., Voss H., Wehland J., Cossart P.;
 RA "Comparative genomics of Listeria species.";
 RT Science 294:849-852(2001).
 RL EMBL; AL596170; CAC97285.1; -.
 DR ListerList; LIN2055; -.
 DR InterPro; IPR002482; LysM.
 DR Pfam; PF01476; LysM; 1.
 DR SMART; SM00257; LysM; 1.
 KM Hypothetical protein; Complete proteome.
 SQ SEQUENCE 243 AA; 25963 MW; 6B2493D143B159D1 CRC64;
 Query Match 42.3%; Score 90; DB 16; Length 243;
 Best Local Similarity 45.5%; Pred. No. 0.13;
 Matches 30; Conservative 3; Mismatches 11; Indels 22; Gaps 2;
 QY 1 AKKTA-KKAKAKKAYKAAEKKA-----AKYKAAAEKAAA 38
 DB 106 AKKAAEEKAAEKAAEKKAADKKSQDEAAKAAAKKQEAEEKAAEKAAA 165
 QY 39 KEAYE 44
 DB 166 DKAAKE 171
 RESULT 6
 ID 001395 PRELIMINARY; PRT; 275 AA.
 AC 001395;
 DT 01-JUL-1997 (TREMBLrel. 04, Created)
 DT 01-JUL-1997 (TREMBLrel. 04, Last sequence update)
 DT 01-OCT-2002 (TREMBLrel. 22, Last annotation update)
 DE Akoneme-associated protein MST101(3).
 GN MST101(3) OR DHMST101.
 OS Drosophila hydei (Fruit fly).
 CC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 CC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 CC Ephydroidea; Drosophilidae; Drosophila.
 CX NCBI_TaxID=7224;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Neesen J., Heinlein U.A.O., Buenemann H.;
 RL Submitted (JUN-1997) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: POSSIBLE STRUCTURAL ROLE IN THE SPERM TAIL (BY
 CC SIMILARITY).
 CC -1- SUBCELLULAR LOCATION: CYTOPLASMIC (BY SIMILARITY).
 CC -1- TISSUE SPECIFICITY: TESTIS (BY SIMILARITY).
 CC -1- DOMAIN: THE PREDOMINANT STRUCTURE IS ALPHA-HELICAL.
 DR EMBL; U85627; AAB51369.1; -.
 DR FLYBase; FBgn0020733; dhyalmer101(3).
 KM Sperm; Repeat; Multigene family.
 FT DOMAIN 64 255
 FT 13 X 16 AA APPROXIMATE TANDEM REPEATS OF
 FT X-[KQ]-K-C-[AE]-E-X-A-[X]-K-X-X-X-X-
 FT [AE]-X.
 SQ SEQUENCE 275 AA; 30436 MW; 76BAA7B2ADDF32C CRC64;

Query Match 41.5%; Score 88.5; DB 5; Length 275;
 Best Local Similarity 61.0%; Pred. No. 0.2;
 Matches 25; Conservative 1; Mismatches 12; Indels 3; Gaps 1;
 QY 5 AKKAAEKAKKAYKAAEKKA-----AAKYKAAAEKAAEKAA 42
 DB 63 AKKCAEAKKEKAAEKKAADKKSQDEAAKAAAKKQEAEEKAAEKAAA 103
 RESULT 7
 ID 09WMX1 PRELIMINARY; PRT; 372 AA.
 AC 09WMX1;
 DT 01-NOV-1999 (TREMBLrel. 12, Created)
 DT 01-NOV-1999 (TREMBLrel. 12, Last sequence update)
 DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
 DE TOLA protein.
 GN TOLA.
 CC Pseudomonas putida.
 CC Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;
 CC Pseudomonadaceae; Pseudomonas.
 CX NCBI_TaxID=303;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=mt-2;
 RX MEDLINE=96198174; PubMed=8626299;
 RA Rodriguez-Herva J.U., Ramos-Gonzalez M.I., Ramos J.;
 RT "The Pseudomonas putida peptidoglycan-associated outer membrane
 RT lipoprotein (PAL) is involved in maintenance of the integrity of the
 RT cell envelope.";
 RT J. Bacteriol. 178:1699-1706(1996).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=mt-2;
 RA Ramos-Gonzalez I.;
 RL Submitted (JUN-1995) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=mt-2;
 RA Rodriguez-Herva J.U.;
 RL Submitted (JUL-1999) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RP SEQUENCE FROM N.A.
 RC STRAIN=mt-2;
 RX MEDLINE=96422022; PubMed=8824639;
 RA Rodriguez-Herva J.U., Ramos J.;
 RT "Characterization of an OprL null mutant of Pseudomonas putida.";
 RL J. Bacteriol. 178:5836-5840(1996).
 DR EMBL; X74218; CAB50780.1; -.
 DR InterPro; IPR005819; Histone_H5.
 DR InterPro; IPR006260; TonB_C.
 DR PRINTS; PRO0624; HISTONEH5.
 DR TIGRPMs; TIGR01352; tonB_Cterm; 1.
 SQ SEQUENCE 372 AA; 40133 MW; 87F49785SC3C0BC CRC64;
 Query Match 41.5%; Score 88.5; DB 2; Length 372;
 Best Local Similarity 39.7%; Pred. No. 0.27;
 Matches 27; Conservative 8; Mismatches 10; Indels 23; Gaps 1;
 QY 1 AKKTA-KKAKAKKAYKAAEKKA-----AKKAA 37
 DB 121 AEDAKAAEAKKAAEKKAADKKSQDEAAKAAAKKQEAEEKAAEKAAA 180
 QY 38 AKAAAYE 45
 DB 181 AEAAXKA 188
 RESULT 8
 ID 09ASJ6 PRELIMINARY; PRT; 899 AA.
 AC 09ASJ6;
 SQ SEQUENCE 275 AA; 30436 MW; 76BAA7B2ADDF32C CRC64;

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DT 01-JUN-2001 (TREMBlrel. 17, Created)
DT 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
DE DNA topoisomerase (EC 5.99.1.2).
GN CC2451.
OS Caulobacter crescentus.
OC Bacteria; Proteobacteria; Alphaproteobacteria; Caulobacteriales;
OC Caulobacteraceae; Caulobacter.
OX NCBI_TaxID=155892;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 19089 / CB15;
RX MEDLINE=21173698; PubMed=11259647;
RA Nieman W.C., Feldblum T.V., Laub M.T., Paulsen I.T., Nelson K.E.,
RA Eisen J., Heidelberg J.F., Alley M.R.K., Ohta N., Maddock J.R.,
RA Porocka I., Nelson W.C., Newton A., Stephens C., Phadke N.D., Ely B.,
RA DeBoy R.T., Dodson R.J., Durkin A.S., Gwin M.L., Haft D.H.,
RA Kolonay J.F., Smit J., Craven M.B., Knouti H., Shetty J., Berry K.,
RA Utecherback T., Tran K., Wolf A., Vamathevan J., Ermolaeva M., White O.,
RA Salzberg S.L., Venter J.C., Shapiro L., Fraser C.M.;
RA Proc. Natl. Acad. Sci. U.S.A. 98:4136-4141(2001).
RT "Complete genome sequence of Caulobacter crescentus."
CC -1- FUNCTION: THE REACTION CATALYZED BY TOPOISOMERASE LEADS TO THE
CC -1- CONVERSION OF ONE TOPOLOGICAL ISOMER OF DNA TO ANOTHER (BY
CC SIMILARITY).
CC -1- CATALYTIC ACTIVITY: ATP-INDEPENDENT BREAKAGE OF SINGLE-STRANDED
CC DNA, FOLLOWED BY PASSAGE AND REJOINING.
CC -1- MISCELLANEOUS: WHEN A TOPOISOMERASE TRANSIENTLY BREAKS A DNA
CC BACKBONE BOND, IT SIMULTANEOUSLY FORMS A PROTEIN-DNA LINK, IN
CC WHICH A TYROSYL OXYGEN IN THE ENZYME IS JOINED TO A DNA PHOSPHORUS
CC AT ONE END OF THE ENZYME-SEVERED DNA STRAND (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO PROKARYOTIC TYPE I/III TOPOISOMERASE
CC FAMILY.
CC DR EMBL: AE005914; AK24422.1; -.
CC DR HSSP: P06612; 1ECL.
CC DR TIGR: CC2451; -.
CC DR InterPro: IPR003601; DNATopI_ATP_bind.
CC DR InterPro: IPR003602; DNATopI_DNA_bind.
CC DR InterPro: IPR005733; DNA_topI_bact.
CC DR InterPro: IPR000380; DNA_topIsmase.
CC DR InterPro: IPR006171; TopoIim_dom.
CC DR InterPro: IPR006154; TopoIim_sub.
CC DR Pfam: PF01131; TopoIsm_bac; 1.
CC DR Pfam: PF01751; TopoIim; 1.
CC DR Pfam: PF01396; zF-C4_TopoIsm; 1.
CC DR PRINTS: PR00417; PRFISMAESI.
CC DR SMART: SM00437; TOP1AC; 1.
CC DR SMART: SM00436; TOP1BC; 1.
CC DR SMART: SM00493; TOP1IM; 1.
CC DR TIGRfams: TIGR01051; topI_bact; 1.
CC DR PROSITE: PS00396; TOPOISOMERASE_I_PROK; 1.
CC DR DNA-binding: Isomerase; Topoisomerase; Complete proteome.
CC DR KEGG: 899 AA; 97723 MW; 1485DCED0ADAF6A CRC64;
CC SQ SEQUENCE

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OC Bacteria; Proteobacteria; Betaproteobacteria; Burkholderiales;
OC Ralstoniaceae; Ralstonia.
OX NCBI_TaxID=305;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=GMT1000;
RX MEDLINE=21681879; PubMed=11823852;
RA Salanoubat M., Genin S., Attiguenave F., Guzy J., Mangenot S.,
RA Arlat M., Billault A., Brottier P., Camus J.C., Cattolico L.,
RA Chandler M., Choisme N., Claudel-Renard C., Cunne S., Demange N.,
RA Gaudin C., Lavie M., Moisan A., Robert C., Sauvin W., Schlex T.,
RA Signier P., Thebaud P., Whalen M., Winkler P., Levy M.,
RA Weisenbach J., Boucher C.A.;
RT "Genome sequence of the plant pathogen Ralstonia solanacearum."
RT Nature 415:497-502(2002).
RX EMBL: AL646071; CAD16500.1; -.
DR InterPro: IPR005819; Histone_H5.
DR PRINTS: PR00624; HISTONEH5.
KM Complete proteome.
SQ SEQUENCE 200 AA; 19279 MW; D3831B590510272D CRC64;

Query Match
Best Local Similarity 40.6%; Score 86.5; DB 16; Length 200;
Matches 26; Conservative 4; Mismatches 11; Indels 5; Gaps 2;

QY 1 AKKYAKKAKKAKKAKKAA---EAKKAKTEKAAEKAAKAA 41
DB 42 AKKYAKKAKKAKKAKKAAKVVAKKAAKAAKAAKAAKAAKAA 87

RESULT 10
ID Q8P140 PRELIMINARY; PRT; 155 AA.
AC Q8P140;
DT 01-OCT-2002 (TREMBlrel. 22, Created)
DT 01-OCT-2002 (TREMBlrel. 22, Last sequence update)
DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
DE Histone H1.
GN XAC3058.
OS Xanthomonas axonopodis (pv. citri).
OC Bacteria; Proteobacteria; Gammaproteobacteria; Xanthomonadales;
OC Xanthomonadaceae; Xanthomonas.
OX NCBI_TaxID=92829;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=306 / ATCC 13902 / XV 101;
RX MEDLINE=22022145; PubMed=12024217;
RA da Silva A.C.R., Ferro J.A., Reinach F.C., Farah C.S., Furlan L.R.,
RA Quaggio R.B., Monteiro-Vitorello C.B., Van Sluys M.A., Almeida N.F.,
RA Alves L.M.C., do Amaral A.M., Bertolini M.C., Camargo L.E.A.,
RA Camarotte G., Camnava F., Cardozo U., Chamberg F., Clapina L.P.,
RA Ciccarelli R.M.B., Coutinho L.L., Curino-Santos J.R., El-Dorri H.,
RA Faria J.B., Ferreira A.J.S., Ferreira R.C.C., Ferro M.I.T.,
RA Fornighieri E.F., Franco M.C., Greggio C.C., Gruber A.,
RA Katayama A.M., Kishi L.T., Leite R.P., Lemos E.G.M., Lemos M.V.F.,
RA Locali B.C., Machado M.A., Madeira A.M.B.N., Martinez-Rossi N.M.,
RA Martins E.C., Medeiros U., Menck C.F.M., Miyaki C.Y., Moon D.H.,
RA Moreira L.M., Rossi M.T.M., Okura V.K., Oliveira M.C., Oliveira V.R.,
RA Pereira H.A., Naves A., Sena J.A.D., Silva C., de Souza R.F.,
RA Spindola L.A.F., Takita M.A., Tamura R.E., Teixeira E.C., Tezza R.I.D.,
RA Trindade dos Santos M., Truffi D., Tsai S.M., White F.F.,
RA Setubal J.C., Kitajima J.P.;
RT "Comparison of the genomes of two Xanthomonas pathogens with differing
RT host specificities."
RT Nature 417:459-463(2002).
RX EMBL: AE011948; AAM37903.1; -.
DR InterPro: IPR005819; Histone_H5.
DR PRINTS: PR00624; HISTONEH5.
KM Complete proteome.
SQ SEQUENCE 155 AA; 16124 MW; FDB53CEB15FD48 CRC64;

Query Match
Best Local Similarity 40.4%; Score 86; DB 16; Length 155;
Matches 26; Conservative 5; Mismatches 12; Indels 2;

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Matches 26; Conservative 4; Mismatches 14; Indels 6; Gaps 2;

Oy 2 KKYAKKAAK-----AKKAYKAAEAKKA-KTEKAAEKAEEAAEAYEA 45
Db 49 KKTAKKATAKKAVKKTAKKATKTAAKKAVKKTAKKATKTAAKKATKTA 98

RESULT 11

O8FP30 ID Q8FP30 PRELIMINARY; PRT; 168 AA.
AC Q8FP30; 01-MAR-2003 (TREMBlrel. 23, Created)
DT 01-MAR-2003 (TREMBlrel. 23, Last sequence update)
DE 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
GN Putative 30S ribosomal protein S16.
OS Corynebacterium efficiens.
OC Bacteria; Actinobacteriales; Actinobacteridae; Actinomycetales;
OC Corynebacteriaceae; Corynebacteriaceae; Corynebacterium.
OX NCBI_TaxID=152794;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=YS-314 / AJ 12310 / DSM 44549 / JCM 11189;
RA Kawarabayashi Y., Yamazaki J., Hino Y., Kikuchi H., Nakamura Y.,
RA Ikeo K., Suzuki M., Maehima J., Itoh T., Yamagishi A., Nishio Y.,
RA Usuda Y., Sugimoto S.;
RT "The entire genomic sequence of Corynebacterium efficiens YS-314.";
RL Submitted (MAY-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AP005220; BAC18770.1; -
KW Ribosomal protein; Complete proteome.
SQ SEQUENCE 168 AA; 18228 MW; 443CE68AA2321C01 CRC64;

Query Match 40.4%; Score 86; DB 16; Length 168;
Best Local Similarity 65.7%; Pred. No. 0.22;
Matches 23; Conservative 3; Mismatches 7; Indels 2; Gaps 1;

Oy 7 KAKAEKAKKAAKAEKAAKAEKAAKAEKAAKAE 41
Db 120 EAITEKKKAREKAEKAEKAA--EKAAEKAATAEA 152

RESULT 12

O93946 ID O93946 PRELIMINARY; PRT; 212 AA.
AC O93946; 01-MAY-1999 (TREMBlrel. 10, Created)
DT 01-MAY-1999 (TREMBlrel. 10, Last sequence update)
DE 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
GN CTX2p (Fragment).
OS Candida albicans (Yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; mitosporic Saccharomycetales; Candida.
OX NCBI_TaxID=5476;
RN [1]
RP SEQUENCE FROM N.A.
RA Kaiser B., Kunkel W., Saluz H.P., Munder T.;
RT "Identification of Candida albicans protein domains with
transcriptional activating properties in Saccharomyces cerevisiae.";
RL Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.
DR EMBL; AJ006637; CA07165.1; -
FT NON TER 1
SQ SEQUENCE 212 AA; 24231 MW; 10C2122E9554A387 CRC64;

Query Match 40.4%; Score 86; DB 3; Length 212;
Best Local Similarity 48.9%; Pred. No. 0.28;
Matches 23; Conservative 10; Mismatches 12; Indels 2; Gaps 1;

Oy 1 AKKAKKA--KAEKAKKAYKAAEAKKAAKAEKAAEKAEEAAEAYEA 45
Db 70 AKKAEKAEKAEKAEKAEKAEKAEKAEKAEKAEKAEKAEKAEKAEKAE 116

RESULT 13

O8W044 ID O8W044 PRELIMINARY; PRT; 98 AA.
AC O8W044; 01-MAR-2002 (TREMBlrel. 20, Created)
DT 01-MAR-2002 (TREMBlrel. 20, Last sequence update)
DE 01-MAR-2002 (TREMBlrel. 23, Last annotation update)
GN LNP18.
OS Leishmania major.
OC Eukaryota; Euklenozoa; Kinetoplastida; Trypanosomatidae; Leishmania.
OX NCBI_TaxID=5664;
RN [1]
RP SEQUENCE FROM N.A.
RA Tzortzis N., Papageorgiou F.T., Tzinia A.K., Soteriadou K.P.;
RT "Identification and characterization of a novel Leishmania gene
encoding for a putative histone H1-like transcription factor.";
RL Submitted (JUN-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AJ237814; CAD21431.1; -
DR InterPro; IPR005819; Histone_H5.
DR PRINTS; PR00624; HISTONEH5.
KW Nuclear protein.
SQ SEQUENCE 98 AA; 9999 MW; 0A4AB93089D6C261 CRC64;

Query Match 39.9%; Score 85; DB 5; Length 98;
Best Local Similarity 55.0%; Pred. No. 0.17;
Matches 22; Conservative 3; Mismatches 15; Indels 0; Gaps 0;

Oy 1 AKKYAKKAEKAEKAYKAAEAKKAAKAEKAAEKAEEAAE 40
Db 59 AKKPAKKVAKKPAKKAKKPAKKPAKKPAKKAKKAAKAAKAAK 98

RESULT 14
O8T9R3 ID O8T9R3 PRELIMINARY; PRT; 111 AA.
AC O8T9R3; 01-JUN-2002 (TREMBlrel. 21, Created)
DT 01-JUN-2002 (TREMBlrel. 21, Last sequence update)
DE 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
GN Leishmania infantum.
OS Leishmania infantum.
OC Eukaryota; Euklenozoa; Kinetoplastida; Trypanosomatidae; Leishmania.
OX NCBI_TaxID=5671;
RN [1]
RP SEQUENCE FROM N.A.
RA Papageorgiou F., Soteriadou K.;
RT "Identification of a Leishmania infantum gene encoding for an histone
H1-like nuclear protein.";
RL Submitted (JUN-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF469106; AAL76335.1; -
KW Nuclear protein.
SQ SEQUENCE 111 AA; 11162 MW; 16168F3B54960B83 CRC64;

Query Match 39.9%; Score 85; DB 5; Length 111;
Best Local Similarity 55.0%; Pred. No. 0.19;
Matches 22; Conservative 3; Mismatches 15; Indels 0; Gaps 0;

Oy 1 AKKYAKKAEKAEKAYKAAEAKKAAKAEKAAEKAEEAAE 40
Db 72 AKKPAKKVAKKPAKKAKKPAKKPAKKPAKKAKKAAKAAKAAK 111

RESULT 15

O45370 ID O45370 PRELIMINARY; PRT; 182 AA.
AC O45370; 01-NOV-1996 (TREMBlrel. 01, Created)
DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)
DE 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
GN Histone H1.
OS Bordetella pertussis.

OC Bacteria; Proteobacteria; Betaproteobacteria; Burkholderiales;
 OC Alcaligenaceae; Bordetella.
 OX NCBI_TaxID=520;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BPM28;
 RX MEDLINE=95319329; PubMed=7596289;
 RA Scariato V., Arico B., Goyard S., Ricci S., Manetti R., Prugnola A.,
 RA Manetti R., Polverino-De-Laureto P., Ullmann A., Rappuoli R.;
 RT "A novel chromatin-forming histone H1 homologue is encoded by a
 RT dispensable and growth-regulated gene in Bordetella pertussis.";
 RL Mol. Microbiol. 15:871-881(1995).
 DR EMBL; L37438; AAB59120.1;
 SQ SEQUENCE 182 AA; 18252 MW; 9A17A397B12B0421 CRC64;

Query Match 39.9%; Score 85; DB 2; Length 182;
 Best Local Similarity 61.4%; Pred. No. 0.3;
 Matches 27; Conservative 4; Mismatches 9; Indels 4; Gaps 3;
 Qy 1 AKKVAKKAAEK--AKKAY-KAAEAKKAAKYEKAAAEKAAAKEA 41
 Db 58 AKKVAKKAAVAKKAAVAKKAAVAKKAAVAKKAAVAKKAAVAKKAA 100

Search completed: January 21, 2004, 09:00:26
 Job time : 21.1169 secs

Rank	Score	Match Length	DB	ID	Description
1	213	100.0	45	21	AAy82572
2	188	64.8	109	21	AAy82577
3	138	63.1	56	21	AAy82573
4	131	59.4	77	21	AAy82575
5	126.5	59.4	86	21	AAy82576
6	120.5	56.6	66	21	AAy82574
7	108	50.7	35	21	AAy82571
8	99.5	46.7	106	11	AAu06446
9	98	46.0	13	AAu08871	High affinity mactrin

Listeria monocytogenes,
Recombinant copoly-
Amino acid sequenc
M. tuberculosis hi
M. tuberculosis h
Novel human diagn
Mycobacterial hepa
Mycobacterial hepa
Mycobacterietal hep
Pseudomonas aerugi
Mycobacterium bovis
C. albicans apoptos
C glutamicum proteo
Tumor neoangiogen
Drosophila melanog
Histone H1 isoform
Human histone HI .5
Histone H1 isoform
Human histone HI .5
Histone H1 isoform
Human histone HI .5
Human histone HI .5
Human histone HI .5
Human histone HI .5
Human histone HI .5
Human histone HI .5
Human histone HI .5
Human histone HI .5
Human histone HI .5
Human ORF protein
Human ovarian anti
Human polypeptide
Human secreted proe
Human secreted prot
Human ribosomal L1
Protein differentie
Human polypeptide

XX (YEND) YEND REC & NEW CO LTD

(YEDA) YEDA RES & DEV CO LTD.

XX W0200018794-A1.
 XX
 XX 06-APR-2000.
 XX
 XX 24-SEP-1999; 99WO-US22402.
 XX
 XX 25-SEP-1998; 98US-0101693.
 XX
 XX (YEDA) YEDA RES & DEV CO LTD.
 XX (TEVA-) TEVA PHARM USA INC.
 XX
 XX Gad A, Lis D;
 XX
 XX WPI; 2000-317499/27.
 XX
 XX Copolymer 1 related polypeptides used as molecular weight markers for
 PT glatiramer acetate and for treatment and prevention of immune diseases
 XX
 XX Claim 10; Page 14; 72pp; English.
 XX
 CC AAY82571 to AAY82577 represent specifically claimed copolymer molecular
 CC weight TV-marker polypeptides from the present invention. The present
 CC invention describes polypeptides (I) for determining the molecular
 CC weight of a copolymer (Cp), which has an identified molecular weight
 CC and an amino acid composition corresponding to the copolymer. The
 CC polypeptides of the invention are used as molecular weight markers for
 CC glatiramer acetate related tetrapolymers. The polypeptides may also be
 CC used for treating and preventing immune diseases in a mammal. Autoimmune
 CC diseases which may be treated include either cell-mediated or
 CC antibody-mediated diseases. Such diseases include arthritic conditions,
 CC demyelinating diseases and inflammatory conditions, e.g. multiple
 CC sclerosis, rheumatoid arthritis, osteoarthritis, autoimmune haemolytic
 CC anaemia, autoimmune oophoritis, autoimmune thyroiditis, autoimmune
 CC uveoretinitis, Crohn's disease, chronic immune thrombocytopenia
 CC purpura, colitis, contact sensitivity disease, diabetes mellitus, Graves
 CC disease, Guillain-Barre's syndrome, Hashimoto's disease, idiopathic
 CC myxoedema, myasthenia gravis, psoriasis, pemphigus vulgaris, or systemic
 CC lupus erythematosus. Mediated-mediated diseases which can be treated
 CC include host-versus-graft disease, graft-versus-host disease, and
 CC delayed-type hypersensitivity. The polypeptides of the invention have
 CC defined molecular weights and physical properties which are analogous to
 CC glatiramer acetate molecules, which makes them ideal for use as
 CC molecular weight markers.
 CC
 XX
 XX Sequence 56 AA;
 SQ
 Query Match 63.1%; Score 134.5; DB 21; Length 56;
 Best Local Similarity 68.4%; Pred. No. 7.2e-08;
 Matches 39; Conservative 0; Mismatches 5; Indels 13; Gaps 4;
 Oy 1 AKKYAKK-----AKAEKA-----KKAYKAAEKK- -AAKYKAAEKKAAYEA 45
 Db 1 AKKYAKKKAAYKAAKKAAYKAAKKAAYKAAKKAAYKAAKKAAYKAAKKAAYEA 56
 RESULT 4
 ID AAY82575 standard; peptide; 77 AA.
 XX
 XX AAY82575;
 XX
 XX 28-JUL-2000 (first entry)
 XX
 XX Copolymer molecular weight TV-marker amino acid sequence SEQ ID NO:5.
 XX
 XX Copolymer; molecular weight marker; TV-marker; immune disease;
 KW glatiramer acetate; autoimmune disease; antiarthritic; neuroprotective;
 KW osteopathic; immunosuppressive; antithyroid; antiinflammatory;
 KW antidiabetic; thyromimetic; haemostatic; antiporiatic; dermatological;
 KW antanaemic; immunosuppressive; demyelinating disease; osteoarthritis;
 KW inflammatory condition; multiple sclerosis; rheumatoid arthritis;
 KW Crohn's disease; chronic immune thrombocytopenia purpura; colitis;

KW diabetes mellitus; Graves disease; Guillain-Barre's syndrome; psoriasis;
 KW Hashimoto's disease; idiopathic myxoedema; myasthenia gravis;
 KW pemphigus vulgaris; systemic lupus erythematosus.
 XX
 XX Unidentified.
 XX
 XX W0200018794-A1.
 XX
 XX 06-APR-2000.
 XX
 XX 24-SEP-1999; 99WO-US22402.
 XX
 XX 25-SEP-1998; 98US-0101693.
 XX
 XX (YEDA) YEDA RES & DEV CO LTD.
 XX (TEVA-) TEVA PHARM USA INC.
 XX
 XX Gad A, Lis D;
 XX
 XX WPI; 2000-317499/27.
 XX
 XX Copolymer 1 related polypeptides used as molecular weight markers for
 PT glatiramer acetate and for treatment and prevention of immune diseases
 XX
 XX Claim 10; Page 14; 72pp; English.
 XX
 CC AAY82571 to AAY82577 represent specifically claimed copolymer molecular
 CC weight TV-marker polypeptides from the present invention. The present
 CC invention describes polypeptides (I) for determining the molecular
 CC weight of a copolymer (Cp), which has an identified molecular weight
 CC and an amino acid composition corresponding to the copolymer. The
 CC polypeptides of the invention are used as molecular weight markers for
 CC glatiramer acetate related tetrapolymers. The polypeptides may also be
 CC used for treating and preventing immune diseases in a mammal. Autoimmune
 CC diseases which may be treated include either cell-mediated or
 CC antibody-mediated diseases. Such diseases include arthritic conditions,
 CC demyelinating diseases and inflammatory conditions, e.g. multiple
 CC sclerosis, rheumatoid arthritis, osteoarthritis, autoimmune haemolytic
 CC anaemia, autoimmune oophoritis, autoimmune thyroiditis, autoimmune
 CC uveoretinitis, Crohn's disease, chronic immune thrombocytopenia
 CC purpura, colitis, contact sensitivity disease, diabetes mellitus, Graves
 CC disease, Guillain-Barre's syndrome, Hashimoto's disease, idiopathic
 CC myxoedema, myasthenia gravis, psoriasis, pemphigus vulgaris, or systemic
 CC lupus erythematosus. Mediated-mediated diseases which can be treated
 CC include host-versus-graft disease, graft-versus-host disease, and
 CC delayed-type hypersensitivity. The polypeptides of the invention have
 CC defined molecular weights and physical properties which are analogous to
 CC glatiramer acetate molecules, which makes them ideal for use as
 CC molecular weight markers.
 CC
 XX
 XX Sequence 77 AA;
 SQ
 Query Match 61.5%; Score 131; DB 21; Length 77;
 Best Local Similarity 50.6%; Pred. No. 2.4e-07;
 Matches 39; Conservative 0; Mismatches 6; Indels 32; Gaps 3;
 Oy 1 AKKYAKK-----AKAEKA-----KKAYKAAEKKAAKYE----- 29
 Db 1 AKKYAKKKAAYKAAKKAAYKAAKKAAYKAAKKAAYKAAKKAAYKAAKKAAYEA 60
 Oy 30 -KAAEKKAAYEA 45
 Db 61 YKAAKKAAYEA 77
 RESULT 5
 ID AAY82576 standard; peptide; 86 AA.
 XX
 XX AAY82576;
 XX
 XX 28-JUL-2000 (first entry)
 XX

XX	Copolymer molecular weight TV-marker amino acid sequence SEQ ID NO:6.
KW	Copolymer; molecular weight marker; TV-marker; immune disease;
KW	glutathione acetate; autoimmune disease; antiarthritic; neuroprotective;
KW	osteoplastic; immunosuppressive; antithyroid; antiinflammatory;
KW	antidiabetic; thyromimetic; haemostatic; antipsoriatic; dermatological;
KW	antihaemic; immunosuppressive; demyelinating disease; osteoarthritis;
KW	inflammatory condition; multiple sclerosis; rheumatoid arthritis;
KW	Crohn's disease; chronic immune thrombocytopenia purpura; colitis;
KW	diabetes mellitus; Graves disease; Guillain-Barre's syndrome; psoriasis;
KW	Hashimoto's disease; idiopathic myxoedema; myasthenia gravis;
KW	pemphigus vulgaris; systemic lupus erythematosus.
OS	unidentified.
XX	
PN	WO200018794-A1.
XX	
PD	06-APR-2000.
XX	
PF	24-SEP-1999; 99WO-US22402.
XX	
PR	25-SEP-1998; 98US-0101693.
XX	
PA	(YEDA) YEDA RES & DEV CO LTD.
PA	(TEVA-) TEVA PHARM USA INC.
XX	
PI	Gad A, Lis D;
DR	WPI; 2000-317499/27.
PT	Copolymer 1 related polypeptides used as molecular weight markers for
PT	glutathione acetate and for treatment and prevention of immune diseases
XX	
XX	Claim 10; Page 14; 72pp; English.

CC	AAV82571.c	AAV82577	represent specifically claimed copolymer molecular
CC	weight TV-marker polypeptides from the present invention. The present		
CC	invention described polypeptides (I) for determining the molecular		
CC	weight of a copolymer (CP), which has an identified molecular weight		
CC	and an amino acid composition corresponding to the copolymer. The		
CC	polypeptides of the invention are used as molecular weight markers for		
CC	glutimer acetate related tetrapolymers. The polypeptides may also be		
CC	used for treating and preventing immune diseases in a mammal. Autoimmune		
CC	diseases which may be treated include either cell-mediated or		
CC	antibody-mediated diseases. Such diseases include arthritic conditions,		
CC	demyelinating diseases and inflammatory conditions, e.g. multiple		
CC	sclerosis, rheumatoid arthritis, osteoarthritis, autoimmune haemolytic		
CC	anaemia, autoimmune oophoritis, autoimmune thyroiditis, autoimmune		
CC	uveoretinitis, Crohn's disease, chronic immune thrombocytopaenia		
CC	purpura, colitis, contact sensitivity disease, diabetes mellitus, Graves		
CC	disease, Guillain-Barre's syndrome, Hashimoto's disease, idiopathic		
CC	myxoedema, myasthenia gravis, psoriasis, pemphigus vulgaris, or systemic		
CC	lupus erythematosus. Mediated-mediated diseases which can be treated		
CC	include host-versus-graft disease, graft-versus-host disease, and		
CC	delayed-type hypersensitivity. The polypeptides of the invention have		
CC	defined molecular weights and physical properties which are analogous to		
CC	glutimer acetate molecules, which makes them ideal for use as		
CC	molecular weight markers.		
XX			
SO	Sequence	86 AA;	
	Query Match	59.4%; Score 126.5; DB 21; Length 86;	
	Best Local Similarity	45.3%; Pred. No. 8.3e-07;	
	Matches	39; Conservative 0; Mismatches 6; Indels 41; Gaps 3	
QY	1 AKTAKK-----AKAEKA-----KAYKAAEKKAATG-----	29	
DB	1 AKYAKKKKKAYAKKKKKAKKAAKAAAYKAAEKKAATGAKYAAKAAEKKEVAABAK	60	
QY	30 -----KAAAKKAAKAAAYEA 45		
DB	61 YKAAAKKAYKAAKAAKAAKAAKAAAYEA 86		

XX	RESULT 6	
XX	AAy82574	
XX	ID	AAy82574 standard; peptide; 66 AA.
XX	AAy82574;	
XX	AC	
XX	DT	
XX	28-JUL-2000	(first entry)
XX	Copolymer molecular weight	TV-marker amino acid sequence SEQ ID NO:4
XX	DE	

KM Copolymer: molecular weight marker; IV-marker: immune disease;
KM glatiramer acetate: autoimmune disease; antiarthritic; neuroprotective;
KM osteopontin; immunosuppressive; antithyroid; antinflammatory;
KM antidiabetic; thymostimetic; haemostatic; antipsoriatic; dermatological;
KM antidiabetic; immunosuppressive; demyelinating disease; osteoarthritis;
KM inflammatory condition; multiple sclerosis; rheumatoid arthritis;
KM Crohn's disease; chronic immune thrombocytopenia purpura; colitis;
KM *Haemobes mellitus*; Graves disease; Guillain-Barre's syndrome; psoriasis
KM Hashimoto's disease; idiopathic myxoedema; myasthenia gravis;
KM *penhphigus vulgaris*; systemic lupus erythematosus.

OS	Unidentified.
XX	
PN	WO200018794-A1.
XX	
PD	06-APR-2000.
XX	
PF	24-SEP-1999; 99WO-US22402.
XX	
PR	25-SEP-1998; 98US-0101693.
XX	
PA	(YEDA) YEDA RES & DEV CO LTD
EA	(TEVA-) TEVA PHARM USA INC.

P1 Gad A, Lis D;
 DR WPI; 2000-317499/27.
 XX
 XX Copolymer 1 related polypeptides used as molecular weight markers for
 PT gastrin acerate and for treatment and prevention of immune diseases
 XX
 XX
 XX Claim 10; Page 14; 72pp; English.

AA18257 to AA18257 represent specifically claimed copolymer molecular weight TV-marker polypeptides from the present invention. The present invention describes polypeptides (1) for determining the molecular weight of a copolymer (CP), which has an identified molecular weight and an amino acid composition corresponding to the copolymer. The polypeptides of the invention are used as molecular weight markers for tetramer acetate related tetrapolymers. The polypeptides may also be used for treating and preventing immune diseases in a mammal. Autoimmune diseases which may be treated include either cell-mediated or antibody-mediated diseases. Such diseases include arthritic conditions, demyelinating diseases and inflammatory conditions, e.g. multiple sclerosis, rheumatoid arthritis, osteoarthritis, autoimmune haemolytic anaemia, autoimmune ophthalmitis, autoimmune thyroiditis, autoimmune uveoretinitis, Crohn's disease, chronic immune thrombocytopenia purpura, colitis, contact sensitivity disease, diabetes mellitus, Graves disease, Guillain-Barre's syndrome, Hashimoto's disease, idiopathic myxoedema, myasthenia gravis, psoriasis, pemphigus vulgaris, or systemic lupus erythematosus. Mediated-mediated diseases which can be treated include host-versus-graft disease, graft-versus-host disease, and delayed-type hypersensitivity. The polypeptides of the invention have defined molecular weights and physical properties which are analogous to tetramer acetate molecules, which makes them ideal for use as molecular weight markers.

Query Match:	56.6%	Score 120.5;	DB 21;	Length 66;
Best Local Similarity:	71.1%	Pred. No. 2.8e-06;		
Matches 32;	Conservative 0;	Mismatches 6;	Indels 7;	Gaps 2

[illegible]

```

50 Sequence 35 AA:
Query Match 50.7%; Score 108; DB 21; Length 35;
Best Local Similarity 64.4%; Pred. No. 3,4e-05;
Matches 29; Conservative 1; Mismatches 5; Indels 10; Gaps 2;

Cy 1 AKKYAKKAKKAKKAYKAAEAKKAYKERYEERAAEKAKEAAVEEA 45
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 1 AKKYAKKEKA-AKKAYK-----KEAKKAAEAAKAAKEAAVEEA 35

RESULT 8
ID AAR06446
AA R06446 standard; protein; 106 AA.
AC AAR06446;
XX
XX
XX 25-MAR-2003 (updated)
DT 03-JAN-1991 (first entry)
XX
XX Recombinant copolymer 1-19, myelin basic protein analogue.
DE
XX
XX Recombinant copolymer 1; COP-1-19; myelin basic protein; MBP;
KW immunological activity; autoimmune encephalomyelitis;
KW multiple sclerosis;
XX
XX Synthetic.
OS
XX
XX EP383620-A.
FN
XX
XX 22-AUG-1990.
PD
XX
XX 16-FEB-1990; 90BP-0301700.
PF
XX
XX 07-FEB-1990; 90US-0473845.
PR 17-FEB-1989; 89US-0312541.
XX
XX (REPK ) REPLIGEN CORP.
PA
XX
XX Cook KS;
PI
XX
XX WPI; 1990-255848/34.
DR N-PSDB; AAQ06446.
XX
XX
XX Producing genes encoding random polymers of aminoacid(s) - for
PT producing recombinant polypeptide(s) with biological and/or
FT immunological activity
XX
XX
XX Disclousure; Fig 12; 25pp; English.
XX
XX
XX To improve the expression of rCOP-1 polypeptides in E. coli, genes
CC coding for rCOP-1-19 were subcloned from pBEV 2.1 to pBG3-2deltaN
CC (deposited: 20-NOV-1984 US4691009, NRRU B-15910), a plasmid used to
CC express Protein A. The resulting plasmids encode fusion proteins
CC consisting of beta-glucuronidase, Protein A, and rCOP-1 sequences.
CC A methionine residue occurs between the Protein A and rCOP-1
CC sequences, originating from the 5' linker sequence, in order that
CC the COP-1 polypeptide may be cleaved from the fusion protein.
CC rCOP-1-19 contains oligonucleotide duplexes encoding the following
CC segments: YKA, AAE, KAA, EKA, KKA, YEA, AKK KAA, and AAA. The
CC N-terminal alanine residue is left behind following CNBr cleavage of the
CC fusion protein.
CC The product prevents or arrests experimental autoimmune
CC encephalomyelitis. They are used to prevent, arrest or control a
CC demyelinating disorder, e.g. multiple sclerosis. They may also
CC be used as additives to hair care products to confer beneficial
CC effects on damaged hair or as supplements for diets deficient in
CC certain amino acids.
CC See also AAQ05664.
CC (Updated on 25-MAR-2003 to correct PA field.)
XX
XX Sequence 106 AA;
50

```


KM	immunological activity; autoimmune encephalomyelitis;
KM	multiple sclerosis;
XX	
XX	Synthetic.
OS	
FN	EP383620-A.
XX	
PD	22-AUG-1990.
XX	
XX	16-FEB-1990; 90EP-0301700.
PF	
XX	
PR	07-FEB-1990; 90US-0473845.
PR	17-FEB-1989; 89US-0312541.
XX	
PA	(REPK) REPLIGEN CORP.
XX	
PI	Cook KS;
XX	
DR	WPI, 1990-255848/34.
DR	N-PSDB; AAQ05664.
XX	
PT	Producing genes encoding random polymers of aminoacid(s) - for
PT	producing recombinant polypeptide(s) with biological and/or
PT	immunological activity
XX	
PS	Disclosure; Fig 11; 25pp; English.
XX	
XX	To improve the expression of rCOP-1 polypeptides in E. coli, genes
CC	coding for rCOP-1-77 were subcloned from pREV 2.1 to PBG3-2delCAN
CC	(deposit: 20-NOV-1984 US4691009, NRRL B-15910), a plasmid used to
CC	express Protein A. The resulting plasmids encode fusion proteins
CC	consisting of beta-glucuronidase, Protein A, and rCOP-1 sequences.
CC	A methionine residue occurs between the Protein A and rCOP-1
CC	sequences, originating from the 5' linker sequence, in order that
CC	the COP-1 polypeptide may be cleaved from the fusion protein.
CC	rCOP-1-77 contains oligonucleotide duplexes encoding the following
CC	segments: YKK, EAG, KAK, AAK, and AAA. The N-terminal alanine residue
CC	is left behind following CNBr cleavage of the fusion protein.
CC	The product prevents or arrests experimental autoimmune
CC	encephalomyelitis. They are used to prevent, arrest or control a
CC	demyelinating disorder, e.g. multiple sclerosis. They may also
CC	be used as additives to hair care products to confer beneficial
CC	effects on damaged hair or as supplements for diets deficient in
CC	certain amino acids.
CC	See also AAQ05665.
XX	(Updated on 25-MAR-2003 to correct PA field.)
XX	
SO	Sequence 154 AA;
XX	
Query Match	44.1%; Score 94; DB 11; Length 154;
Best Local Similarity	60.5%; Pred. No. 0.0052;
Matches	26; Conservative 2; Mismatches 9; Indels 6; Gaps 2;
XX	
YY	2 KCVAKKAKAEKAKAYKQAEKKAKYKAEKAAAEKAAEAYE 44
DB	105 KKYKKKAKAKYKK--KAKEAKA-----KAAAEKAKAEKAYK 141
XX	
RESULT 12	
AAV14928	
ID	AAV14928 standard; protein; 223 AA.
XX	
AC	AAV14928;
XX	
XX	25-OCT-1999 (first entry)
DE	
XX	Amino acid sequence of M. vaccae antigen GV-45.
XX	
KM	Mycobacterium vaccae protein; antigen; T cell activation; cytokine;
KM	dendritic cell maturation; infectious disease; immune disorder; cancer;
KM	respiratory system; mycobacterial infection; allergy; tuberculosis;
KM	leprosy; sarcoidosis; lung cancer; asthma; skin disorder; psoriasis;
KM	dermatitis; eczema; alopecia areata; skin cancer; basal carcinoma;

KW squamous cell carcinoma; melanoma.
 XX
 OS Mycobacterium vaccae.
 XX
 XX WO9932634-A2.
 PN
 XX
 PD 01-JUL-1999.
 XX
 PF 23-DEC-1998; 98WO-NZ00189.
 XX
 XX 04-DEC-1998; 98US-0205426.
 PR 23-DEC-1997; 97US-0996624.
 PR 23-DEC-1997; 97US-0997080.
 PR 23-DEC-1997; 97US-0997362.
 PR 11-JUN-1998; 98US-0095855.
 PR 17-SEP-1998; 98US-0156181.
 XX
 PA (GENE-) GENESIS RES & DEV CORP LTD.
 XX
 PI Prestidge RL, Skinner MA, Tan P, Visser ES, Watson J;
 DR WPI; 1999-430163/36.
 DR N-PSDB; AA211393.
 XX
 PT Enhancing immune response to an antigen
 XX
 PS Claim 1; Page 239; 243pp; English.
 XX
 CC The invention provides heat-killed Mycobacterium vaccae, or recombinant
 CC M. vaccae proteins. The M. vaccae proteins may be employed to activate
 CC T cells and natural killer cells, to stimulate the production of
 CC cytokines, to enhance the expression of co-stimulatory molecules on
 CC dendritic cells and monocytes, and to enhance dendritic cell maturation
 CC and function. The proteins can be expressed by standard recombinant
 CC methodology. Pharmaceutical compositions comprising the proteins or
 CC nucleic acid sequences encoding the proteins can be used for the
 CC treatment, prevention, and detection of disorders including infectious
 CC diseases, immune disorders and cancer. In particular, the compounds and
 CC methods are used for treatment of diseases of the respiratory system,
 CC such as mycobacterial infections, asthma, allergies, tuberculosis,
 CC leprosy, sarcoidosis and lung cancers, and disorders of the skin such as
 CC psoriasis, atopic dermatitis, eczema, allergic contact dermatitis,
 CC alopecia areata, and skin cancers such as basal carcinoma, squamous cell
 CC carcinoma and melanoma.
 CC
 SQ Sequence 223 AA;
 Query Match 42.7%; Score 91; DB 20; Length 223;
 Best Local Similarity 59.6%; Pred. No. 0.016;
 Matches 28; Conservative 5; Mismatches 12; Indels 2; Gaps 2;
 QY 1 AKKVA-KKAKAEKAKKAYCAAEKKAAYEKA-AEKAAKEAYEA 45
 DB 137 AKKATKAPAKKATPAKKAAPAKKATPAKKAAPAKKAPAKKATPA 183

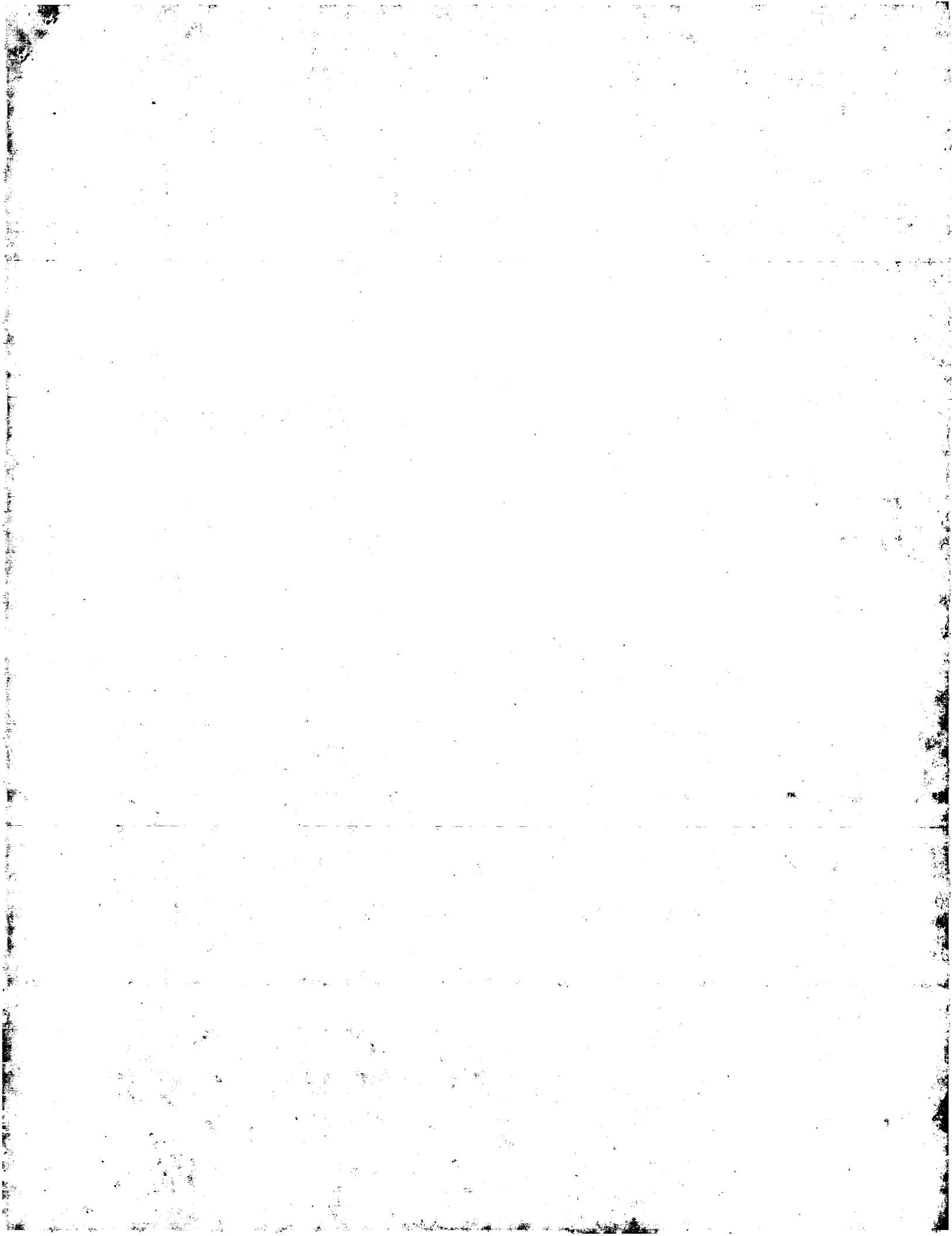
RESULT 13
 ID AAY34055
 ID AAY34055 standard; protein; 214 AA.
 XX
 AC AAY34055;
 XX
 DT 23-NOV-1999 (first entry)
 XX
 DE M. tuberculosis histone H1-like antigen.
 XX
 KW Ulcerative colitis; histone; H1-like antigen; porin antigen;
 KW Bacteroides antigen; inflammatory bowel disease; IBD; PANCA; diagnosis;
 KW perinuclear anti-neutrophil cytoplasmic antibody.
 XX
 OS Mycobacterium tuberculosis.
 XX
 PN WO9945955-A1.

XX
 PD 16-SEP-1999.
 XX
 PF 12-MAR-1999; 99WO-US05492.
 XX
 PR 12-MAR-1998; 98US-0041889.
 XX
 PA (REGC) UNIV CALIFORNIA.
 XX
 PI Braun J, Cohavy O;
 DR WPI; 1999-551215/46.
 PT Use of histone H1, porin or Bacteroides antigens as targets for the
 PT diagnosis, prevention and treatment of ulcerative colitis -
 XX
 PS Claim 2; Fig 11; 134pp; English.
 XX
 CC The invention provides a method for the diagnosis, prevention and
 CC treatment of ulcerative colitis (UC) using histone H1-like antigen, a
 CC porin antigen or a Bacteroides antigen as a target antigen. The novel
 CC method of diagnosing UC in a subject suspected of having inflammatory
 CC bowel disease (IBD) comprises: (1) obtaining a sample from the subject;
 CC (2) contacting the sample with a histone H1-like antigen, or perinuclear
 CC anti-neutrophil cytoplasmic antibody (panCA)-reactive fragment, to form a
 CC complex of the histone H1-like antigen, or the panCA-reactive fragment,
 CC and antibody to the histone H1-like antigen; and (3) detecting the
 CC presence or absence of the complex; where the presence of the complex
 CC indicates that the subject has UC. The panCA-reactive histone H1-like
 CC antigen, porin antigen and Bacteroides antigen are useful in the
 CC diagnosis, prevention and treatment of UC. The methods can also be used
 CC for identifying agents useful for treating UC. The present sequence
 CC represents a M. tuberculosis histone H1-like antigen.
 CC
 SQ Sequence 214 AA;
 Query Match 42.0%; Score 89.5; DB 20; Length 214;
 Best Local Similarity 55.6%; Pred. No. 0.023;
 Matches 25; Conservative 4; Mismatches 13; Indels 3; Gaps 1;
 QY 1 AKKVAKKAKAEKAKKAYCAAEKKAAYEKA-AEKAAKEAYEA 45
 DB 111 AKKAKKAPAKKATPAKKAATPAKKAATPAKKAATPAKKAATPA 152

RESULT 14
 ID AAY57353
 ID AAY57353 standard; Protein; 214 AA.
 XX
 AC AAY57353;
 XX
 DT 13-JUN-2000 (first entry)
 XX
 DE M. tuberculosis histone H1-like protein, 214.
 XX
 KW Ulcerative colitis; inflammatory bowel disease; porin antigen; MB;
 KW PANCA; perinuclear anti-neutrophil cytoplasmic antibody; 214 protein;
 KW histone H1; isoform.
 XX
 OS Mycobacterium tuberculosis.
 XX
 PN US6033864-A.
 XX
 PD 07-MAR-2000.
 XX
 PF 12-MAR-1998; 98US-0041889.
 XX
 PR 12-APR-1996; 96US-0057846.
 PR 11-APR-1997; 97US-0837058.
 XX
 PA (REGC) UNIV CALIFORNIA.
 XX
 PI Cohavy O, Braun J;

XX DR WPI; 2000-255695/22.
 XX
 PT Diagnosing ulcerative colitis or susceptibility, by detecting complex
 PT formation between microbial porin antigen and perinuclear
 PT anti-neutrophil cytoplasmic autoantibodies -
 XX
 PS Example 5; Fig 10; 49pp; English.
 XX
 CC The invention provides a method for diagnosing ulcerative colitis in a
 CC subject suspected of having inflammatory bowel disease. The method
 CC comprises reacting a patient sample with a porin antigen that is
 CC immunologically reactive with pANCA (perinuclear anti-neutrophil
 CC cytoplasmic antibodies) and detecting formation of a Ag-pANCA complex
 CC as indicative of ulcerative colitis. The method is used to diagnose
 CC ulcerative colitis or susceptibility to it. The present sequence
 CC represents a histone H1-like protein of M. tuberculosis, designated 214.
 XX
 SQ Sequence 214 AA;
 Query Match 42.0%; Score 89.5; DB 21; Length 214;
 Best Local Similarity 55.6%; Pred. No. 0.023;
 Matches 25; Conservative 4; Mismatches 13; Indels 3; Gaps 1;
 Oy 1 AKKYAKKAKKAKKAYKAAEAKKAKYKAAAEKAAKAAEAA 45
 Db 111 AKKYAKKAPAKKATPAKAAKATKAPA--RKAATPAKAKATPA 152
 |||||:|||||:|||||:|||||:|||||:|
 RESULT 15
 ABG28693
 ID ABG28693 standard; Protein; 334 AA.
 XX
 AC ABG28693;
 XX
 DT 18-FEB-2002 (first entry)
 XX
 DE Novel human diagnostic protein #28684.
 XX
 KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder.
 XX
 OS Homo sapiens.
 XX
 FN WO200175067-A2.
 XX
 PD 11-OCT-2001.
 XX
 PE 30-MAR-2001; 2001WO-US08631.
 XX
 PR 31-MAR-2000; 2000US-0540217.
 PR 23-AUG-2000; 2000US-0649167.
 XX
 PA (HYSE-) HYSEQ INC.
 XX
 PI Drmanac RT, Liu C, Tang YT;
 XX
 DR WPI; 2001-639362/73.
 DR N-PSDB; AAS92880.
 XX
 PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -
 XX
 PS Claim 20; SEQ ID No 59052; 103pp; English.
 XX
 CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques

CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. ABG00010-ABG30377 represent novel human
 CC diagnostic amino acid sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 XX
 SQ Sequence 334 AA;
 Query Match 40.8%; Score 87; DB 22; Length 334;
 Best Local Similarity 57.4%; Pred. No. 0.067;
 Matches 27; Conservative 6; Mismatches 12; Indels 2; Gaps 2;
 Oy 1 AKKYAKKAKKAKK-KAYKAAEAKKAKYKAAAEKAA-KEAA 45
 Db 214 AKKYATETAEKKAQADKKAABKKAADKKAABKKAATKKA 260
 |||||:|||||:|||||:|||||:|||||:|
 Search completed: January 21, 2004, 08:59:09
 Job time : 22.6234 secs



RESULT 4
US-09-816-989A-5
; Sequence 5, Application US/09816989A
; Patent No. US20020115103A1
; GENERAL INFORMATION:
; APPLICANT: Gad, Alexander

```

QY      30 -----KAAAEKAAAEAAVEA 45
          ||| ||| ||| ||| ||| |||
Db      61 YKAEAAKAYKAEAAKAAAEAAVEA 86

```

RESULT 6


```

US-09-816-989A-4
; Sequence 4, Application US/09816989A
; Patent No. US20020115103A1
; GENERAL INFORMATION:
; APPLICANT: Gad, Alexander
; APPLICANT: Lis, Doris
; TITLE OF INVENTION: COPOLYMER 1 RELATED POLYPEPTIDES FOR USE AS MOLECULAR WEIGHT MARK
; FILE REFERENCE: 2609/60807-A-PCT-US
; CURRENT FILING DATE: 2001-03-23
; PRIOR FILING DATE: 1998-09-25
; PRIOR APPLICATION NUMBER: 60/101,693
; PRIOR FILING DATE: 1999-09-24
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 4
; LENGTH: 66
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic Peptide
US-09-816-989A-4

Query Match          56.6%; Score 120.5; DB 10; Length 66;
Best Local Similarity 71.1%; Pred. No. 2.2e-06;
Matches 32; Conservative 0; Mismatches 6; Indels 7; Gaps 2;

Cy 1 AKKYAKKAKAEKAKYKAAEKKAAYEKAAYEA 45
Db 29 AKKYAKKAKAE-KKEYAAAEK-----YKAAKAAKAAKAAAYEA 66

RESULT 7
US-09-816-989A-1
; Sequence 1, Application US/09816989A
; Patent No. US20020115103A1
; GENERAL INFORMATION:
; APPLICANT: Gad, Alexander
; APPLICANT: Lis, Doris
; TITLE OF INVENTION: COPOLYMER 1 RELATED POLYPEPTIDES FOR USE AS MOLECULAR WEIGHT MARK
; FILE REFERENCE: 2609/60807-A-PCT-US
; CURRENT FILING DATE: 2001-03-23
; PRIOR FILING DATE: 1998-09-25
; PRIOR APPLICATION NUMBER: 60/101,693
; PRIOR FILING DATE: 1999-09-24
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 1
; LENGTH: 35
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic Peptide
US-09-816-989A-1

Query Match          50.7%; Score 108; DB 10; Length 35;
Best Local Similarity 64.4%; Pred. No. 2.5e-05;
Matches 29; Conservative 1; Mismatches 5; Indels 10; Gaps 2;

Cy 1 AKKYAKKAKAEKAKYKAAEKKAAYEKAAYEA 45
Db 1 AKKYAKKAKAEKAKYKAAEKKAAYEKAAYEA 35

RESULT 8
US-10-205-979-52
; Sequence 52, Application US/10205979
; Publication No. US20030147861A1

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```

; GENERAL INFORMATION:
; APPLICANT: Watson, James D.
; APPLICANT: Tan, Paul L. J.
; APPLICANT: Abernethy, Nevin
; TITLE OF INVENTION: Compounds and Methods for the Modulation
; FILE REFERENCE: 11000.1063U
; CURRENT FILING DATE: 2002-07-25
; PRIOR FILING DATE: 2001-07-26
; NUMBER OF SEQ ID NOS: 52
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 52
; LENGTH: 223
; TYPE: PRT
; ORGANISM: Mycobacterium vaccae
US-10-205-979-52

Query Match          42.7%; Score 91; DB 12; Length 223;
Best Local Similarity 59.6%; Pred. No. 0.013;
Matches 28; Conservative 5; Mismatches 12; Indels 2; Gaps 2;

Cy 1 AKKYA-KKAYAKKAKYKAAEKKAAYEKA-AEKAAAEKAAAYEA 45
Db 137 AKKAATTAAPAKKATTAAKKAAPAKGATPAKKAAPAKKAATPA 183

RESULT 9
US-10-051-643-201
; Sequence 201, Application US/10051643
; Publication No. US20020197265A1
; GENERAL INFORMATION:
; APPLICANT: Watson, James D.
; APPLICANT: Tan, Paul L. J.
; TITLE OF INVENTION: Methods and Compounds for the Treatment
; FILE REFERENCE: 11000.1008C2
; CURRENT FILING DATE: 2002-01-18
; PRIOR FILING DATE: 1997-12-23
; PRIOR APPLICATION NUMBER: US 08/996,624
; PRIOR FILING DATE: 1998-09-17
; NUMBER OF SEQ ID NOS: 208
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 201
; LENGTH: 223
; TYPE: PRT
; ORGANISM: Mycobacterium vaccae
US-10-051-643-201

Query Match          42.7%; Score 91; DB 14; Length 223;
Best Local Similarity 59.6%; Pred. No. 0.013;
Matches 28; Conservative 5; Mismatches 12; Indels 2; Gaps 2;

Cy 1 AKKYA-KKAKAEKAKYKAAEKKAAYEKA-AEKAAAEKAAAYEA 45
Db 137 AKKAATTAAPAKKATTAAKKAAPAKGATPAKKAAPAKKAATPA 183

RESULT 10
US-10-229-567-27
; Sequence 27, Application US/10229567
; Publication No. US20030092080A1
; GENERAL INFORMATION:
; APPLICANT: Braun, Jonathan
; APPLICANT: Chavvy, Ofer
; TITLE OF INVENTION: Diagnosis, Prevention and Treatment of
; Ulcerative Colitis, and Clinical Subtypes Thereof, Using
; Microbial UC PANCA antigens
; NUMBER OF SEQUENCES: 41

```


TYPE: PRT
ORGANISM: Corynebacterium glutamicum
US-09-738-626-5751

Query Match 38.5%; Score 82; DB 10; Length 165;
Best Local Similarity 62.9%; Pred. No. 0.087;
Matches 22; Conservative 4; Mismatches 7; Indels 2; Gaps 1;

QY 7 KAKAEKAKKAYKAAEAKKAAKYEKAAEKAKEA 41
DB 120 EAITEKKKKAREDEKAEKA--EKAAAEKAAAS 152

RESULT 14
US-10-156-761-12370
Sequence 12370, Application US/10156761
Publication No. US20030119018A1
GENERAL INFORMATION:
APPLICANT: OMURA, SATOSHI
APPLICANT: IKEDA, HARUO
APPLICANT: ISHIKAWA, JUN
APPLICANT: HORIKAWA, HIROSHI
APPLICANT: SHIBA, TADAYOSHI
APPLICANT: SAKAKI, YOSHIYUKI
APPLICANT: HATTORI, MASAHIRA
TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
FILE REFERENCE: 249-262
CURRENT APPLICATION NUMBER: US/10/156,761
CURRENT FILING DATE: 2002-05-29
PRIOR APPLICATION NUMBER: JP 2001-204089
PRIOR FILING DATE: 2001-05-30
PRIOR APPLICATION NUMBER: JP 2001-272697
PRIOR FILING DATE: 2001-08-02
NUMBER OF SEQ ID NOS: 15109
SEQ ID NO 12370
LENGTH: 272
TYPE: PRT
ORGANISM: Streptomyces avermitilis
US-10-156-761-12370

Query Match 36.6%; Score 78; DB 15; Length 272;
Best Local Similarity 46.3%; Pred. No. 0.41;
Matches 19; Conservative 7; Mismatches 15; Indels 0; Gaps 0;

QY 1 AKKYAKKAKAEKAKKAYKAAEKKAAKYEKAAEKAKEA 41
DB 90 AAKAKQAKSDLADAKKKAETKKAARRAAERAAASRSA 130

RESULT 15
US-09-820-843A-24
Sequence 24, Application US/09820843A
Publication No. US20030039963A1
GENERAL INFORMATION:
APPLICANT: Council of Scientific and Industrial Research
TITLE OF INVENTION: A COMPUTATIONAL METHOD FOR THE IDENTIFICATION OF CANDIDATE PROTEI
FILE REFERENCE: Q63915
CURRENT APPLICATION NUMBER: US/09/820,843A
CURRENT FILING DATE: 2001-03-30
NUMBER OF SEQ ID NOS: 118
SOFTWARE: Patencin version 3.0
SEQ ID NO 24
LENGTH: 309
TYPE: PRT
ORGANISM: Pseudomonas aeruginosa
FEATURE:
NAME/KEY: misc.feature
OTHER INFORMATION: polyhydroyalcanoate synthesis protein Phaf
NAME/KEY: misc.feature
OTHER INFORMATION: g1|9951352
US-09-820-843A-24

Query Match 36.6%; Score 78; DB 11; Length 309;
Best Local Similarity 54.8%; Pred. No. 0.47;
Matches 23; Conservative 0; Mismatches 19; Indels 0; Gaps 0;

QY 1 AKKYAKKAKAEKAKKAYKAAEKKAAKYEKAAEKAKEA 42
DB 142 AKAAKPAKPAKPAKPAKTAAPKPAKPAKPAKPAKPA 183

Search completed: January 21, 2004, 09:10:08
Job time : 16.3636 secs


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; GENERAL INFORMATION:
; APPLICANT: Yeda Research and Development Co., Ltd.
; TITLE OF INVENTION: GLATIRAMER ACETATE MOLECULAR WEIGHT MARKERS

```

```

QY      1 AKKYA-KKAAERAKKAAYKAAEAKKAAYEKAA-AEKAAKEAYAERA 45
        |||||:::||::|||::|||::|||::|||::|||::|||::|||::|||
Db      137 AKGAATGAPAKKATAAKKAAPAKKATPAKKAAAPAKKAAAPAKKAATKA 183

RESULT 9
US-09-205-426-201
Sequence 201, Application US/09205426
Patent No. 6406704
GENERAL INFORMATION:
APPLICANT: Watson, James D.
TITLE OF INVENTION: Compounds and Methods for Treatment and
FILE REFERENCE: 11000.1002c4
CURRENT APPLICATION NUMBER: US/09/205,426
EARLIER FILING DATE: 1998-12-04
EARLIER APPLICATION NUMBER: 09/095,855
EARLIER FILING DATE: 1998-06-11
EARLIER APPLICATION NUMBER: 08/997,362
EARLIER FILING DATE: 1997-12-23
EARLIER APPLICATION NUMBER: 08/873,970
EARLIER FILING DATE: 1997-06-12
EARLIER APPLICATION NUMBER: 08/705,347
EARLIER FILING DATE: 1996-08-29
NUMBER OF SEQ ID NOS: 208
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 201
LENGTH: 223
TYPE: PR1
ORGANISM: Mycobacterium vaccae
US-09-205-426-201

Query Match          42.7%; Score 91; DB 4; Length 223;
Best Local Similarity 59.6%; Pred. No. 0.0062;
Matches 28; Conservative 5; Mismatches 12; Indels 2; Gaps 2;

QY      1 AKKYA-KKAAERAKKAAYKAAEAKKAAYEKAA-AEKAAKEAYAERA 45
        |||||:::||::|||::|||::|||::|||::|||::|||::|||
Db      137 AKGAATGAPAKKATAAKKAAPAKKATPAKKAAAPAKKAAAPAKKAATKA 183

RESULT 10
US-09-041-889-27
Sequence 27, Application US/09041889
Patent No. 6033864
GENERAL INFORMATION:
APPLICANT: Braun, Jonathan
APPLICANT: Cohavy, Ofer
TITLE OF INVENTION: Diagnosis, Prevention and Treatment of
TITLE OF INVENTION: Ulcerative Colitis, and Clinical Subtypes Thereof, Using
TITLE OF INVENTION: Microbial UC pAMCA antigens
NUMBER OF SEQUENCES: 41
CORRESPONDENCE ADDRESS:
ADDRESSEE: Campbell & Flores LLP
STREET: 4370 La Jolla Village Drive, Suite 700
CITY: San Diego
STATE: California
COUNTRY: USA
ZIP: 92122
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/041,889
FILING DATE:

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```

; LENGTH: 214 amino acid
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-09-417-264-27

```

Qy 1 AKCYAKKAKAEKAKKAYKAAEAKKAACTYERKAAAEKAAKEAA 42
157 AKAAAKPPAKPPAKPPAKPTAAAKPPAKPPAKKAAAKPPAKPPAA 198

RESULT 14
US-09-107-532A-5094

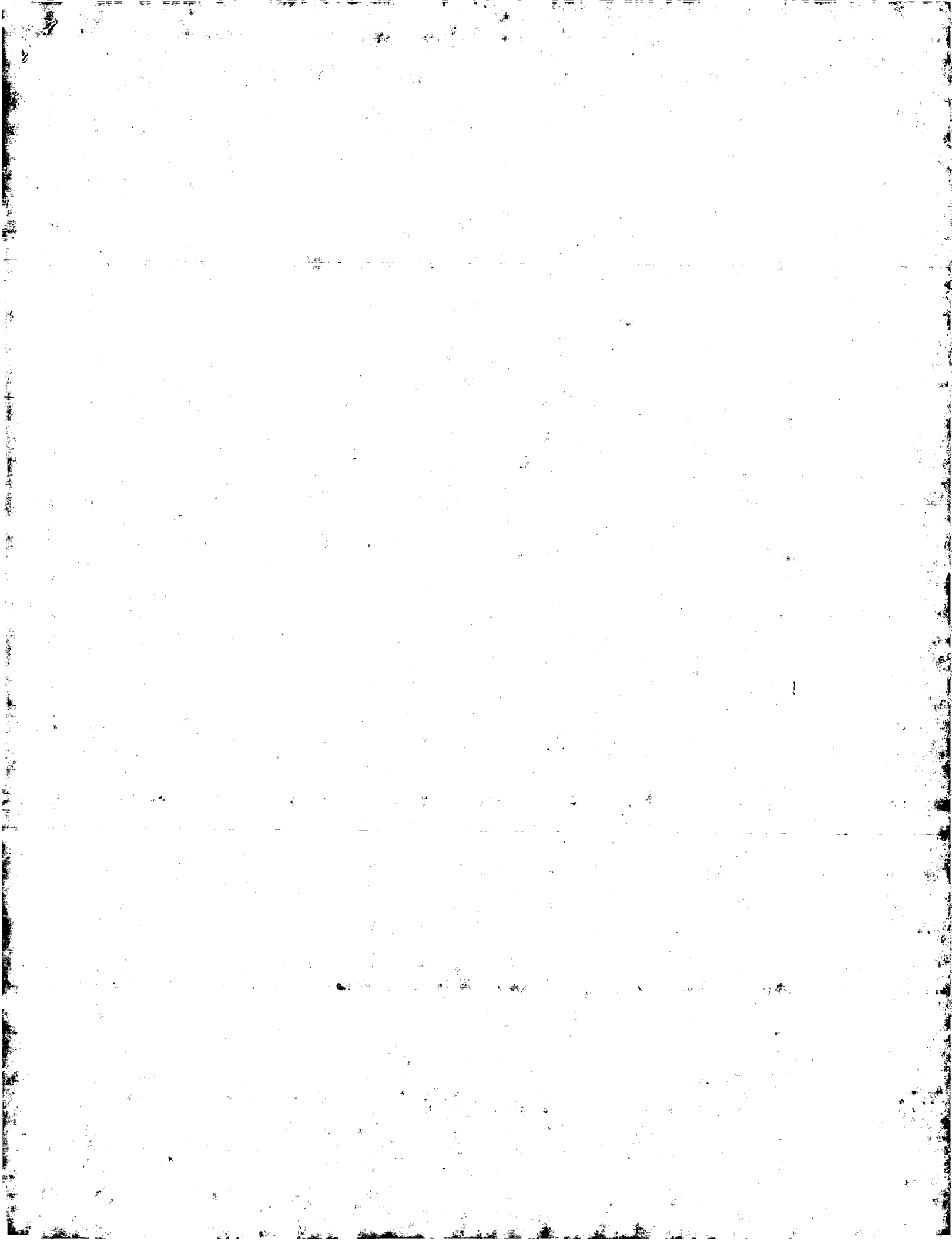
RESULT 2

RESULT 1

[illegible]

RESULT 15
S19113
cgcr-4 protein - Chlamydomonas reinhardtii (fragment)
C|Species: Chlamydomonas reinhardtii
C|Date: 13-Jan-1995 #sequence_revision 13-Jan-1995 #text_change 21-Jul-2000
C|Accession: S19113; S14466
R|Wakarchuk, W.W.; Mueller, F.W.; Beck, C.F.
Plant Mol. Biol. 18, 143-146, 1992
A|Title: Two GC-rich DNA elements of Chlamydomonas reinhardtii with complex arrangements
A|Reference number: S19113; MUID:92119224; PMID:1731966
A|Accession: S19113
A|Status: Preliminary
A|Molecule type: DNA
A|Residues: 1-265 <WAK>
C|Cross-references: EMBL:X17208; NID:g18136; PIDN:CAA35080.1; PID:g18137
C|Genetics:
A|Gene: cgcr-4

Search completed: January 21, 2004, 09:01:40
Job time : 20.8182 secs




```

RA Ltloubee R.; central domain interacts with Escherichia coli porins." ;
RT "TolA central domain interacts with Escherichia coli porins." ;
RL EMOO J. 15:6408-6415(1996).
[6]
RN X-RAY CRYSTALLOGRAPHY (1.85 ANGSTROMS) OF 298-421.
RP MEDLINE=99332679; PubMed=10404600;
RX Lubkowski J., Hennecke F., Plueckthun A., Wlodawer A.;
RT "Flamentouse phase infection: crystal structure of gfp in complex
with its coreceptor, the C-terminal domain of TolA." ;
RL Structure 7:711-722(1999).
CC -1- FUNCTION: INVOLVED IN THE TONB-INDEPENDENT UPTAKE OF GROUP A
COLICINS (COLICINS A, B1, E2, E3, AND K). NECESSARY FOR THE
COINCING TO REACH THEIR RESPECTIVE TARGETS AFTER INITIAL
BINDING TO THE BACTERIA. ALSO INVOLVED IN THE TRANSLLOCATION
OF BACTERIOPHAGE DNA.
CC -1- SUBUNIT: INTERACTS, VIA DOMAIN II, WITH PORINS OMPC, OMPC, PHOE
AND LAMB.
CC -1- SUBCELLULAR LOCATION: Type II membrane protein. Inner membrane.
-----
CC CC This SWISS-PROT entry is copyright. It is produced through a collaboration
between the Swiss Institute of Bioinformatics and the EMBL outstation
at the European Bioinformatics Institute. There are no restrictions on
its use by non-profit institutions as long as its content is in no way
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entities requires a license agreement (See http://www.isb-sib.ch/announce/
or send an email to license@isb-sib.ch).
CC -----
DR EMBL; M28232; AAA24683.1; -.
DR EMBL; AE000177; AAC73833.1; -.
DR EMBL; D90713; BAA35405.1; -.
DR PIR; JY0057; JY0057.
DR PDB; 1TOL; 20-MAY-99.
DR Ecocode; EG11007; tolA.
KW Transport; Protein transport; Bacteriocin transport; Transmembrane;
Repeat; Inner membrane; 3D-structure; Complete proteome.
FT DOMAIN 1 13 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 14 34 POTENTIAL.
FT DOMAIN 35 421 PERITPLASMIC (POTENTIAL).
FT DOMAIN 48 310- DOMAIN II (ALPHA-HELICAL).
FT DOMAIN 311 421 DOMAIN III. (FUNCTIONAL).
FT DOMAIN 224 278 10 X TANDEM REPEATS OF [ED]-K(1,2)-
A(2,4).
FT FT
FT HELIX 335 349
FT TURN 350 351
FT TURN 353 354
FT HELIX 355 358
FT TURN 359 360
FT STRAND 363 369
FT TURN 371 372
FT STRAND 375 383
FT HELIX 385 397
FT HELIX 406 412
FT TURN 413 414
FT STRAND 416 421
SO SEQUENCE .421 AA; 43156 MW; 8B2F52B4B97C655E CRC64;
Query Match 36.5%; Score 189.5; DB 1; Length 421;
Best Local Similarity 53.0%; Pred. No. 4.6e-06;
Matches 61; Conservative 11; Mismatches 32; Indels 11; Gaps 5
QY 1 AKTKAKAE---KAVAKKAKEKKVAKKEAVAYOAEEKKAKAEKKYAK-EAAK 55
DY 120 AEEAKQAELKOKQAEEEAARAAAADAKKVAEADAVA--AEBAKKTAADDAKKKAEEAAK 177
QY 56 AKKEAY-QAEAKKYAKAAKAEKGEFAAAEAK--AEAAYKAPAAATPAAEAA 106
DB 178 AAEEKQAEEAAAAALKKKAEEAAEAABARRKKAATEAEAKKAEKKAAAEKA 232
RESULT 2
HI_LYTP1 STANDARD; PRT; 210 AA.
ID HI_LYTP1
DC P06144;
```

```

DE 01-JAN-1998 (Rel. 06, Created)
DT 01-JAN-1998 (Rel. 06, Last sequence update)
DT 15-JUN-1999 (Rel. 38, Last annotation update)
DE Late histone H1.
OS Lytechinus pictus (Painted sea urchin).
OC Eukaryota; Metazoa; Echinodermata; Eleutherozoa; Echinozoa;
OC Echinoida; Euechinoidea; Echinacea; Temnopleurozoa; Toxopneustidae;
OC Lytechinus.
RX NCBI_TaxID=7653;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Embryo;
RX MEDLINE=87040778; PubMed=3022245;
RA Knowles J.A., Childs G.J.;
RT "Comparison of the late H1 histone genes of the sea urchins
RL Lytechinus pictus and Strongylocentrotus purpuratus."
CC Nucleic Acids Res. 14:8121-8133(1986).
CC -1- FUNCTION: HISTONES H1 ARE NECESSARY FOR THE CONDENSATION OF
CC NUCLEOSOME CHAINS INTO HIGHER ORDER STRUCTURES.
CC -1- SUBCELLULAR LOCATION: Nuclear.
CC -1- SIMILARITY: BELONGS TO THE HISTONE H1/H5 FAMILY.
CC -----
CC This SWISS-PROT entry is copyrighted. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its use by
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (see http://www.isb.ch/announce).
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; X04488; CAA28177.1; -.
DR PIR; A25550; A25550.
DR HSSP; P02259; IHST.
DR InterPro; IPR005818; Histone_H1/H5.
DR InterPro; IPR005819; Histone_H5.
DR InterPro; IPR003216; Linkerhist_N.
DR Pfam; PF00538; linker histone; I.
DR PRINTS; PR00624; HISTONH5.
DR PRODOM; PD000373; Linkerhist_N; 1.
DR SMART; SM00526; H1S; 1.
DR Chromosomal protein, Nuclear protein, DNA-binding, Multigene family.
KW SEQUENCE 210 AA; 21746 MW; 08C38F6494007E2 CRC64;
SQ
Query Match 34.9%; Score 181; DB 1; Length 210;
Best Local Similarity 50.0%; Pred. No. 9.3e-06;
Matches 58; Conservative 8; Mismatches 36; Indels 14; Gaps 4
QY 2 KKVAKKAEKVAKKAKAKKEGVAAYKAEKAYKAAE-----AKKAKAEAKKYAKAEAKAK 57
DB 96 KTEAQKA-PAAKAKAKLAANKKEGKKAATTKARKETLAANKAKAKKAKKAYKKPAAPAK 154
QY 58 KEAYKAEAKKYAKAKAKAEKKEVAAAEAKKA-----EAAKAYKAAEAKKAAEAA 106
DB 155 KPAAKKAKKPAAK--KAKKRPAKKPAKKAKKAPPAKKAAPAKKAAKKPAKKA 208
RESULT 3
TOLA_PSEAB
ID TOLA_PSEAB STANDARD; PRT; 347 AA.
AC P50600.
DT 01-OCT-1996 (Rel. 34, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE TOLA protein.
GN TOLA OR PA0971.
OS Pseudomonas aeruginosa.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;
OC Pseudomonadaceae; Pseudomonas.
OX NCBI_TaxID=287;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=PAO;
RX MEDLINE=97113525; PubMed=8955385;

```

RA Dennis J.J., Lafontaine E.R., Sokol P.A.:
 RT "Identification and characterization of the tolDRA genes of
 RL Pseudomonas aeruginosa.";
 RN J. Bacteriol. 178:7059-7068 (1996).
 (2)
 RA REVISIONS TO N-TERMINUS.
 RN Duan K., Sokol P.A.;
 RL Submitted (AUG-1999) to the EMBL/Genbank/DBJ databases.
 (3)
 RP SEQUENCE FROM N.A.
 RC STRAIN=ATCC 15692 / PAOI;
 RX MEDLINE=20437337; PubMed=10984043;
 RA Stover C.K., Pham X.-Q.T., Erwin A.L., Mizoguchi S.D., Warren P.,
 RA Hickey M.T., Brinkman F.S.L., Hutnagle W.O., Kowalik D.J., Lagrou M.,
 RA Gader R.L., Goltz L., Tolentino E., Westbrock-Wadman S., Yuan Y.,
 RA Brody L.L., Coulter S.N., Folger K.R., Kas A., Laidig K., Lam R.M.,
 RA Smith K.A., Spencer D.H., Wong G.K.-S., Wu Z., Paulsen I.T.,
 RA Reizer J., Sater M.H., Hancock R.E.W., Lory S., Olson M.V.;
 RT "Complete genome sequence of Pseudomonas aeruginosa PAOI, an
 RT opportunistic pathogen.";
 RL Nature 406:959-964 (2000).
 CC -1- FUNCTION: INVOLVED IN THE TONB-INDEPENDENT UPTAKE OF PROTEINS
 CC (BY SIMILARITY).
 CC -1- SUBCELLULAR LOCATION: Type II membrane protein. Inner membrane
 CC (Potential).
 CC -----
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 CC -----
 CC EMBL: U39558; AAC4660.2; -;
 DR EMBL: AB004530; AAG04360.1; -;
 DR PIR: B83525; B83525.
 DR InterPro: IPR006260; TonB_C.
 DR TIGFAPS: TIG01352; tonB_Cterm; 1.
 KW Transport; Protein transport; Transmembrane; Repeat; Inner membrane;
 KW Complete proteome.
 FT DOMAIN 1 16 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 17 37 POTENTIAL.
 FT DOMAIN 38 347 PERIPLASMIC (POTENTIAL).
 FT DOMAIN 209 216 POLY-ALA.
 SQ SEQUENCE 347 AA; 37935 MW; EBD4B04A095945 CRC64;
 Query Match 34.5%; Score 179; DB 1; Length 347;
 Best Local Similarity 49.1%; Pred. No. 1.9e-05;
 Matches 54; Conservative 17; Mismatches 23; Indels 10; Gaps 4;
 QY 6 KKAERAKVAKAKAEK-----KAYAKKAKVAKAEKKAQAKAKAKVAKAEKAKAE 59
 DB 99 QKLEGGQVAAAKAAKKADEKRAKAEQAQAAKKADEKKAQAEKKAQKQADIAKKR 158
 QY 60 AKKAAKKYAAKAKKEKKEVAAAEEKAEAAKAYAEAAKAAKAAKAAVEA 109
 DB 159 A-EDEAKK--KAAEDAKKK--AAEDAKKKAABEAKKKAABEAKKAAVEA 204
 RESULT 4
 MST1_DROHY STANDARD; PRT; 344 AA.
 AC Q08695;
 DT 01-FEB-1995 (Rel. 31, Created)
 DT 01-FEB-1995 (Rel. 31, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Axoneme-associated protein met101(1).
 GN MST101(1).
 OS Drosophila hydei (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydriidae; Drosophilidae; Drosophila.

OX NCBI_TaxID=7224;
 RN [1]
 RP SEQUENCE FROM N.A., AND CHARACTERIZATION.
 RC TISSUE=Testis;
 RX MEDLINE=94200512; PubMed=8150205;
 RA Neesen J., Buemann H., Heinlein U.A.;
 RT "The Drosophila hydei gene Dhmet101(1) encodes a testis-specific,
 RT repetitive, axoneme-associated protein with differential abundance in
 RT Y chromosomal deletion mutant flies.";
 RL Dev. Biol. 162:414-425 (1994).
 CC -1- FUNCTION: POSSIBLE STRUCTURAL ROLE IN THE SPERM TAIL. IT IS
 CC ASSOCIATED WITH AXONEMAL STRUCTURES.
 CC -1- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
 CC -1- TISSUE SPECIFICITY: TESTIS. LOCATED IN SPERMATOCTES AND
 CC SPERMATID BUNDLES.
 CC -1- DOMAIN: THE PREDOMINANT STRUCTURE IS ALPHA-HELICAL.
 CC -1- POLYMORPHISM: THE NUMBER OF REPEATS VARIES BETWEEN STRAINS.
 CC -----
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 CC -----
 CC EMBL: X73480; CAAS1875.1; -;
 DR PIR: S34153; S34153.
 DR FlyBase: FBgn011816; Dhyl\met101(1).
 KW Sperm; Repeat; Multigene family.
 FT DOMAIN 58 337
 FT 19 X 16 AA APPROXIMATE TANDEM REPEATS OF
 SQ SEQUENCE 344 AA; 37793 MW; 24C65D2510387E2A CRC64;
 Query Match 33.8%; Score 175.5; DB 1; Length 344;
 Best Local Similarity 49.2%; Pred. No. 3.1e-05;
 Matches 58; Conservative 8; Mismatches 35; Indels 17; Gaps 5;
 QY 1 AKKYAKKAEK---AYAKKAAKAEKKAAYAKKAAE-----AKKAAKAEKCY 49
 DB 65 AKKEKAAEKKKCAEAAKKEKAEKKAQAKKCAEAAKKEKKAQAEKKEKAEKCK 124
 QY 50 AKKAAKAKKAEKKAQAKKAEKKEVAAAEEK--AEAAKYKKAATAAKAEAA 106
 DB 125 CAEAAKKEKA--AEKKCAEAAKKEK--AAEKKCAEAAKKEKAEKKAQAEAA 177
 RESULT 5
 MST2_DROHY STANDARD; PRT; 1391 AA.
 AC Q08696;
 DT 01-FEB-1995 (Rel. 31, Created)
 DT 01-FEB-1995 (Rel. 31, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Axoneme-associated protein met101(2).
 GN MST101(2).
 OS Drosophila hydei (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydriidae; Drosophilidae; Drosophila.
 OX NCBI_TaxID=7224;
 RN [1]
 RP SEQUENCE FROM N.A., AND CHARACTERIZATION.
 RX MEDLINE=9504538; PubMed=7957199;
 RA Neesen J., Padmanabhan S., Buemann H.;
 RT "Tandemly arranged repeats of a novel highly charged 16-amino-acid
 RT motif representing the major component of the sperm-tail-specific
 RT axoneme-associated protein family Dhmet101 form extended
 RT alpha-helical rods within the extremely elongated spermatzoa of
 RT Drosophila hydei.";
 RL Eur. J. Biochem. 225:1089-1095 (1994).
 CC -1- FUNCTION: POSSIBLE STRUCTURAL ROLE IN THE SPERM TAIL.
 CC -1- SUBCELLULAR LOCATION: Cytoplasmic.

[illegible]

Query Match	Best Local Similarity	Score	DB 1	Length	248	Matches	52	Conservative	9	Mismatches	41	Indels	11	Gaps	2
Qy	2	KKVAKKAEKAVAKKAKAEKKA	EKA	KKVAKKAEKKA	EKA	KKVAKKAEKKA	EKA	KKVAKKAEKKA	EKA	KKVAKKAEKKA	EKA	KKVAKKAEKKA	EKA	KKVAKKAEKKA	EKA
Db	120	KKAKKTSAAAKAKKAKKAA	AKKAKKAKKAKKAA	AKKAKKAKKAKKAA	AKKAKKAKKAKKAA	AKKAKKAKKAKKAA	AKKAKKAKKAKKAA	AKKAKKAKKAKKAA	AKKAKKAKKAKKAA	AKKAKKAKKAKKAA	AKKAKKAKKAKKAA	AKKAKKAKKAKKAA	AKKAKKAKKAKKAA	AKKAKKAKKAKKAA	AKKAKKAKKAKKAA
Qy	62	KA--EAKKAK--	-----	AAKKEKKE	YAAAEAKK	KA	BAKAKAY	AE	AKA	AAK	103				
Db	180	KA	AKKAKKAKKAKKAKKAKK	AKKAKKAKKAKKAKKAKK	AKKAKKAKKAKKAKKAKK	AKKAKKAKKAKKAKKAKK	AKKAKKAKKAKKAKKAKK	AKKAKKAKKAKKAKKAKK	AKKAKKAKKAKKAKKAKK	AKKAKKAKKAKKAKKAKK	AKKAKKAKKAKKAKKAKK	AKKAKKAKKAKKAKKAKK	AKKAKKAKKAKKAKKAKK	AKKAKKAKKAKKAKKAKK	AKKAKKAKKAKKAKKAKK
RESULT 7															
H1B_STRPU															
ID	H1B_STRPU	STANDARD;		PRT;											
AC	P15869;														
DT	01-APR-1990	(Rel. 14, Created)													
DT	01-APR-1990	(Rel. 14, Last sequence update)													
DT	15-JUL-1999	(Rel. 38, Last annotation update)													
DE	Histone H1-beta, late embryonic.														
OS	Strongylocentrotus purpuratus	(Purple sea urchin).													
OC	Eukaryota; Metazoa; Echinodermata; Eleutherozoa; Echinozoa;														
OC	Echinoidae; Euechinoidae; Echinacea; Echinoida; Strongylocentrotidae;														
OC	Strongylocentrotus.														
OX	NCBI_TaxID=7668;														
RN	[1]														
RP	SEQUENCE FROM N.A.														
RX	MEDLINE=88246461; PubMed=2837660;														
RA	Lai Z.-C., Childs G.,														
RT	"Characterization of the structure and transcriptional patterns of														
RT	the gene encoding the late histone subtype H1-beta of the sea urchin														
RL	Strongylocentrotus purpuratus."														
RL	Mol. Cell. Biol. 8:1842-1844(1988).														
CC	- FUNCTION: HISTONES H1 ARE NECESSARY FOR THE CONDENSATION OF														
CC	NUCLEOSOME CHAINS INTO HIGHER ORDER STRUCTURES.														
CC	- SUBCELLULAR LOCATION: Nuclear.														
CC	- SIMILARITY: BELONGS TO THE HISTONE H1/H5 FAMILY.														
CC															
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CC	use by non-profit institutions as long as its content is in no way														
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CC	entities requires a license agreement (see http://www.isb-sib.ch/announce/														
CC	or send an email to license@isb-sib.ch).														
CC															
DR	EMBL; M20314; AAA30052.1; -.														
DR	PIR; A28100; A28100.														
DR	HSSP; P02259; IHST.														
DR	InterPro; IPR005818; Histone_H1/H5.			</											

Query Match 32.7%; Score 169.5; DB 1; Length 211;
 Beest Local Similarity 48.7%; Pred. No. 5.1e-05;
 Matches 55; Conservative 8; Mismatches 41; Indels 9; Gaps 4;

QY 3 KYAKKAE-KAYAKKA--KAAEKKAAYAKKAAKAAK--KKAAYAKKAYAKKAAK---55
 DB 88 KLGKKKKKSPDAQKAPDAKAKAKAAKKKKAAKKAAKKEKLAAKKSKTTTKYK 147
 QY 56 --AKKAYKAEKKYAKKAAKKEKYYAAAEEKKAPAAKAYKAAKAAKAAKANA 106
 DB 148 KPAKAKKAPKAAKAAKAPKAAKAAKAAKAPAKKAAKAAKAPAKKAAKAAKAAKVA 200

RESULT 8
 FAU_DROME STANDARD; PRT; 668 AA.
 ID FAU DROME
 AC Q9VFX3; Q9VFX1; Q9VFX2; Q9YOF9;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE Anoxia upregulated protein.
 GN FAU OR CG6544.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.
 OC NCBI_TaxID=7227;
 RP SEQUENCE FROM N.A.
 RC STRAIN=Canton-S; TISSUE=Head;
 RX MEDLINE=99097004; PubMed=9878744;
 RA Ma E., Xu T., Haddad G.G.;
 RT "Gene regulation by O2 deprivation: an anoxia-regulated novel gene in
 Drosophila melanogaster.";
 RL Brain Res. Mol. Brain Res. 63:217-224(1999).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Berkley;
 RX MEDLINE=20196006; PubMed=10731132;
 RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
 RA Amaratunga P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,
 RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
 RA Sutton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
 RA Brandon R.C., Rogers Y.-H.C., Blazer R.G., Champagne M., Pfeiffer B.D.,
 RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L.G.,
 RA Abril J.F., Agbayani A., An H.-J., Andrews-Pfannkoch C., Baldwin D.,
 RA Ballew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
 RA Beeson K.Y., Bessie P.V., Bertone P.P., Bhandari D., Bolshakov S.,
 RA Borokova D., Botchan M.R., Bouck J., Brokstein P., Brotlier P.,
 RA Butris K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra I.,
 RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
 RA de Pablos B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
 RA Dodson K.J., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
 RA Durbin K.J., Evangelista C.C., Ferraz C., Ferrieres S., Fleischmann W.,
 RA Foster C., Gabrielian A.E., Garg N.S., Gelbart W.M., Glasser K.,
 RA Glodex A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
 RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
 RA Houston D., Houston K.A., Howland T.J., Wei M.-H., Ibegwam C.,
 RA Jajala M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
 RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
 RA Lasoko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
 RA Liu X., Maltsev B., McIntosh T.C., McLeod M.P., McPherson D.,
 RA Mouton S.M., Moy M., Murphy B., Murphy C., Morris J., Mohtrefi A.,
 RA Nelson D.R., Nelson K.A., Nixon K.A., Nusskern D.R., Pacle J.M.,
 RA Palazolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
 RA Relarte K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
 RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
 RA Spier E., Spirdaly A.C., Stapleton M., Strong R., Sun E.,
 RA Svendsen R., Tecor C., Turner R., Venter E., Wang A.H., Wang X.,
 RA Wang Z.-Y., Weissman D.A., Weinstein G.M., Weissbach J.,
 RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,

RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
 RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
 RT "The genome sequence of Drosophila melanogaster.";
 RL Science 287:2185-2195(2000).
 RN [3]
 RP REVISIONS, AND ALTERNATIVE SPLICING.
 RC STRAIN=Berkley;
 RX MEDLINE=22426069; PubMed=12537572;
 RA Misra S., Crosby M.A., Mungall C.J., Matthews B.B., Campbell K.S.,
 RA Hradecky P., Huang Y., Kaminker J.S., Millburn G.H., Prochuk S.E.,
 RA Smith C.D., Tupy J.L., Whitfield E.J., Bayraktaroglu L., Bertman B.P.,
 RA Betencourt B.R., Celniker S.E., de Grey A.D.N.J., Drysdale R.A.,
 RA Harris N.L., Richter J., Russo S., Schroeder A.J., Shu S.Q.,
 RA Stapleton M., Yamada C., Ashburner M., Gelbart W.M., Rubin G.M.,
 RA Lewis S.E.;
 RT "Annotation of the Drosophila melanogaster euchromatic genome: a
 RT systematic review.";
 RL Genome Biol. 3:RESEARCH0083.1-RESEARCH0083.22(2002).
 RN [4]
 RP SEQUENCE FROM N.A. (ISOFORM E).
 RC STRAIN=Berkley; TISSUE=Head;
 RX MEDLINE=22426066; PubMed=12537569;
 RA Stapleton M., Carlson J.W., Brokstein P., Yu C., Champagne M.,
 RA George R.A., Guarnin H., Krommiller B., Pacle J.M., Park S., Wan K.H.,
 RA Rubin G.M., Celniker S.E.;
 RT "A Drosophila full-length cDNA resource.";
 RL Genome Biol. 3:RESEARCH0080.1-RESEARCH0080.8(2002).
 CC -1- FUNCTION: Plays an important role in the regulation of tissue
 CC responsiveness to oxygen deprivation.
 CC -1- ALTERNATIVE PRODUCTS:
 CC Event-Alternative splicing. Named isoforms=5;
 CC Comment=Experimental confirmation may be lacking for some
 CC isoforms;
 CC Name=A;
 CC IsoId=Q9VFX3-1; Sequence=Displayed;
 CC Name=B;
 CC IsoId=Q9VFX3-2; Sequence=VSP_004048, VSP_004049;
 CC Name=C;
 CC IsoId=Q9VFX3-3; Sequence=VSP_004046, VSP_004047;
 CC Name=D;
 CC IsoId=Q9VFX3-4; Sequence=VSP_004050, VSP_004051;
 CC Name=E;
 CC IsoId=Q9VFX3-5; Sequence=VSP_004052;
 CC -1- TISSUE SPECIFICITY: Concentrated in lamina neurons, first optic
 CC lobe neurons and cortical neurons of central brain.
 CC -1- INDUCTION: By anoxia.
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 CC -----
 CC EMBL; AF154418; AAD38397.1; -
 CC EMBL; AE003688; AAF54549.2; -
 CC EMBL; AE003688; AAF54550.1; -
 CC EMBL; AE003688; AAF54551.1; -
 CC EMBL; AE003688; AAF54552.1; -
 CC EMBL; AY060997; AAL28545.1; -
 CC EMBL; AY119569; AAM50223.1; -
 CC FLYBase; FBgn0020439; fau.
 CC GO; GO:0006979; P:response to oxidative stress; IDA.
 KW Alternative splicing.
 FT DOMAIN 88 143
 FT THR-GLU-
 FT ALA-GLU-RICH.
 FT IYSVEKTRVKSIPYISGVSRRVVGATRVTSPIRV
 FT TSPARVSRVHSPPVAVVTRRVISSPERTVSTTPS
 FT TVSPSYLSTYTSYTVFTSYTV -> TRPDLCTDRGS
 FT HRSRASDYSYTSKSVSKSVSDSNPSPHSPTSPSTV
 FT EKTSSRGSGSYNSTERTSTTGAGPGSYVSTTSGMLPG

CC	-1- SIMILARITY: BELONGS TO THE HISTONE H1/H5 FAMILY.
CC	-----
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CC	or send an email to license@isb-sib.ch .
CC	-----
DR	EMBL; M17019; AAAA8789.1; .-
DR	PIR; B28456; B28456.
DR	PDB; 1GHC; 31-AUG-94.
DR	InterPro; IPRO05818; Histone H1/H5.
DR	InterPro; IPRO05819; Histone H5.
DR	Pfam; PF00538; linker_histone_1.
DR	PRINTS; PRO0624; HISTONH5.
DR	SMART; SM00526; H1s; 1.
KW	Chromosomal protein; Nuclear protein; DNA-binding; Multigene family;
KW	Acetylation; 3D-structure.
FT	INIT MET 0 0
FT	MOD RES 1 1 ACETYLATION (BY SIMILARITY).
FT	DOMAIN 40 113 GLOBULAR.
FT	TURN 44 54
FT	HELIx 54 54
FT	TURN 62 63
FT	HELIx 66 68
FT	HELIx 81 87
FT	TURN 88 90
FT	HELIx 91 94
FT	TURN 95 95
FT	TURN 104 105
SQ	SEQUENCE 224 AA; 22397 MW; D3D057CB97865CAF CRC64;
Query Match	31.4%; Score 163; DB 1; Length 224;
Best Local Similarity	43.8%; Pred. No. 0.0001;
Matches 49;	Conservative 13; Mismatches 42; Indels 8; Gaps 2;
Cy	1 AKKKAKKAEEKAYAKAAKEKK-----AVAKKEAKYVAAEKKKAKAAEKKVAKEA 53
Dd	112 SKRGGEVKERAPKKASPAKPPEAKPPAAPAAKKPKKAAVAVKKSPKKAKPPASATKS 171
Cy	54 AKAKEAVYKAEEAKKYAKAAVEKKEVAA-AAEAKKAEAKAKYKAEEAAVAAKE 104
Dd	172 AKSKKVTTKAVKPKKAAVAAKSPAKAKAVKPRPAAPKPAKAPRAAKAKKAAK 223
RESULT 12	
ID	H1_ONCMY STANDARD; PRT; 206 AA.
AC	PO6350;
DT	01-JAN-1988 (Rel. 06, Created)
DT	01-JAN-1988 (Rel. 06, Last sequence update)
DT	15-JUL-1999 (Rel. 38, Last annotation update)
DE	Histone H1.
OS	Oncorhynchus mykiss (Rainbow trout) (Salmo gairdneri).
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC	Actinopterygii; Neopterygii; Teleostei; Euteleostei;
OC	Protacanthopterygii; Salmoniformes; Salmonidae; Oncorhynchus.
OX	NCBI_Taxid=8022;
RN	[1]
RP	SEQUENCE FROM N.A.
EX	MEDLINE=65264847; PubMed=6443128.
RA	Mequitta U., Connor W., Winktein R.J., Dixon G.H.;
RT	"An H1 histone gene from rainbow trout (Salmo gairdneri).";
RL	J. Mol. Evol. 21:209-219(1985).
CC	-1- FUNCTION: HISTONES H1 ARE NECESSARY FOR THE CONDENSATION OF NUCLEOSOME CHAINS INTO HIGHER ORDER STRUCTURES.
CC	-1- SUBCELLULAR LOCATION: Nuclear.
CC	-1- SIMILARITY: BELONGS TO THE HISTONE H1/H5 FAMILY.

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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: X02624; CAB37646.1; -.
CC PIR: A02584; HSTRIR.
CC HSSP: P08287; IGRC.
CC InterPro: IPR005818; Histone H1/H5.
CC InterPro: IPR005819; Histone H5.
CC Pfam: PF00538; linker histone; 1.
CC PRINTS: PR00624; HISTONEH5.
CC SMART: SM00526; H15; 1.
CC Chromosomal protein; Nuclear protein; DNA-binding; Multigene family;
CC Acetylation.
CC INIT MET 0 0 ACETYLATION (BY SIMILARITY).
CC MOD RES 1 1 GLOBULAR.
CC DOMAIN 27 100
CC SEQUENCE 206 AA; 20672 MW; 72C440798066716C CRC64;
CC
CC Query Match 30.9%; Score 160.5; DB 1; Length 206;
CC Best Local Similarity 47.7%; Pred. No. 0.00019;
CC Matches 52; Conservative 9; Mismatches 35; Indels 13; Gaps 4;
CC
QY 1 AKKVAK-----KAEKVAKAKAKAKKVAKKAKEKVAKAAEAKKAKAKAKKVAKEAK 55
DB 105 AKKPAKAAAPKAKKAAKPAKAAKPAKAAKPAKAAKPAKAAKPAKAAKPAKAAKPAK 164
QY 56 AKKEAYKAEKKAAYKAKAKAEKKEVAAAEKKAEEAAKAAKAAKAAKAAK 104
DB 165 VKKPA--AAAK---KAAKSPK--ATKAAKPAKAAKPAKAAKAAKAAK 205

RESULT 13
H15_HUMAN STANDARD; PRT; 225 AA.
AC P16701; Q14529;
DT 01-AUG-1990 (Rel. 15, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Histone H1.5 (Histone H1a).
GN HISTH1B OR H1P5.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OC NCBI_TaxID=9606;
OX (1)
RN RP SEQUENCE.
RC TISSUE=SPLEEN;
RX MEDLINE=90130391; PubMed=2613692;
RA "Human spleen histone H1. Isolation and amino acid sequences of three
RT minor variants, H1a, H1c, and H1d."
RL J. Biochem. 106:844-857(1989).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=97183654; PubMed=9011620;
RA Albig W., Meerjans T., Doenacke D.;
RT "Characterization of the H1.5 gene completes the set of human H1
RL subtype genes."
RL Gene 184:141-148(1997).
RN [3]
RP SEQUENCE FROM N.A.
RX MEDLINE=22296985; PubMed=12408966;
RA Marxluff W.F., Gongidi P., Woods K.R., Jin J., Malais L.J.;
RT "The human and mouse replication-dependent histone genes."
RL Genomics 80:487-498(2002).
RN [4]
RP SEQUENCE FROM N.A.

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RA Wild A.;
RL Submitted (DEC-1997) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: HISTONES H1 ARE NECESSARY FOR THE CONDENSATION OF
CC NUCLEOSOME CHAINS INTO HIGHER ORDER STRUCTURES.
CC -!- SUBCELLULAR LOCATION: Nuclear.
CC -!- SIMILARITY: BELONGS TO THE HISTONE H1/H5 FAMILY.
CC -----
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CC -----
CC EMBL: X83509; CAA58498.1; -.
CC EMBL: AF531304; AAN06704.1; -.
CC PIR: Z98744; CAB11421.1; -.
CC PIR: S51660; S51660.
CC HSSP: P08287; IGRC.
CC Genew; HGNC:4719; HISTH1B.
CC MIM: 142711; -.
CC GO: GO:0005718; C:nucleosome; NAS.
CC GO: GO:0003677; F:DNA binding activity; NAS.
CC GO: GO:0007001; P:chromosome organization and biogenesis (see. . .; NAS.
CC GO: GO:0006334; P:nucleosome assembly; NAS.
CC InterPro: IPR005818; Histone H1/H5.
CC InterPro: IPR005819; Histone H5.
CC Pfam: PF00538; linker histone; 1.
CC PRINTS: PR00624; HISTONEH5.
CC SMART: SM00526; H15; 1.
CC Chromosomal protein; Nuclear protein; DNA-binding; Multigene family;
CC Acetylation.
CC INIT MET 0 0
CC MOD RES 1 1 ACETYLATION.
CC CONFLICT 215 217 MISSING (IN REF. 1).
CC SEQUENCE 225 AA; 22449 MW; 26CD4A1E5D463CDA CRC64;
CC
CC Query Match 30.9%; Score 160.5; DB 1; Length 225;
CC Best Local Similarity 46.8%; Pred. No. 0.0002;
CC Matches 51; Conservative 7; Mismatches 44; Indels 7; Gaps 2;
CC
QY 1 AKKVAKAEKKAAYKAKAKAEKKAAYKAAKAAKAAKAAKAAK 55
DB 118 AKPKAKKGAAPKAKKPAKATPKK--AKKAAKAAKAAKVKTPKPAKAAKAAKAAKSPK 175
QY 56 AKKEAYKAEKKAAYKAKAKAEKKEVAAAEKKAEEAAKAAKAAKAAKAAK 104
DB 176 AKAAKPKKATKSPKPAKPAKPAKPAKPAKPAKPAKPAKPAKPAKPAKPAK 224

RESULT 14
DBH_MYCSM STANDARD; PRT; 208 AA.
AC Q9ZEC5;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE DNA-binding protein HU homolog (Histone-like protein) (H1p).
GN HUP OR H1P.
OS Mycobacterium smegmatis.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Corynebacteriaceae; Mycobacteriaceae; Mycobacterium.
OX NCBI_TaxID=1772;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 700084 / mc(2)155;
RX MEDLINE=99110209; PubMed=9894918;
RA Lee B.H., Murugaou-Oel B., Dick T.;
RT "Upregulation of a histone-like protein in dormant Mycobacterium
RL smegmatis."
RL Mol. Gen. Genet. 260:475-479(1998).
CC -!- FUNCTION: THIS PROTEIN BELONGS TO THE HISTONE LIKE FAMILY OF

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DR PDB; 1GHC; 31-AUG-94.
DR InterPro; IPR005818; Histone_H1/H5.
DR InterPro; IPR005819; Histone_H5.
DR Pfam; PF00538; linker_histone_1.
DR PRINTS; PR00624; HISTONEH5.
DR SMART; SM00526; H15; 1.
DR Chromosomal protein; Nuclear protein; DNA-binding; Multigene family;
KW 3D-structure.
FT INIT_MET 0
FT DOMAIN 37 110 GLOBULAR.
SQ SEQUENCE 218 AA; 21672 MW; CB9724BPF14654A6 CRC64;

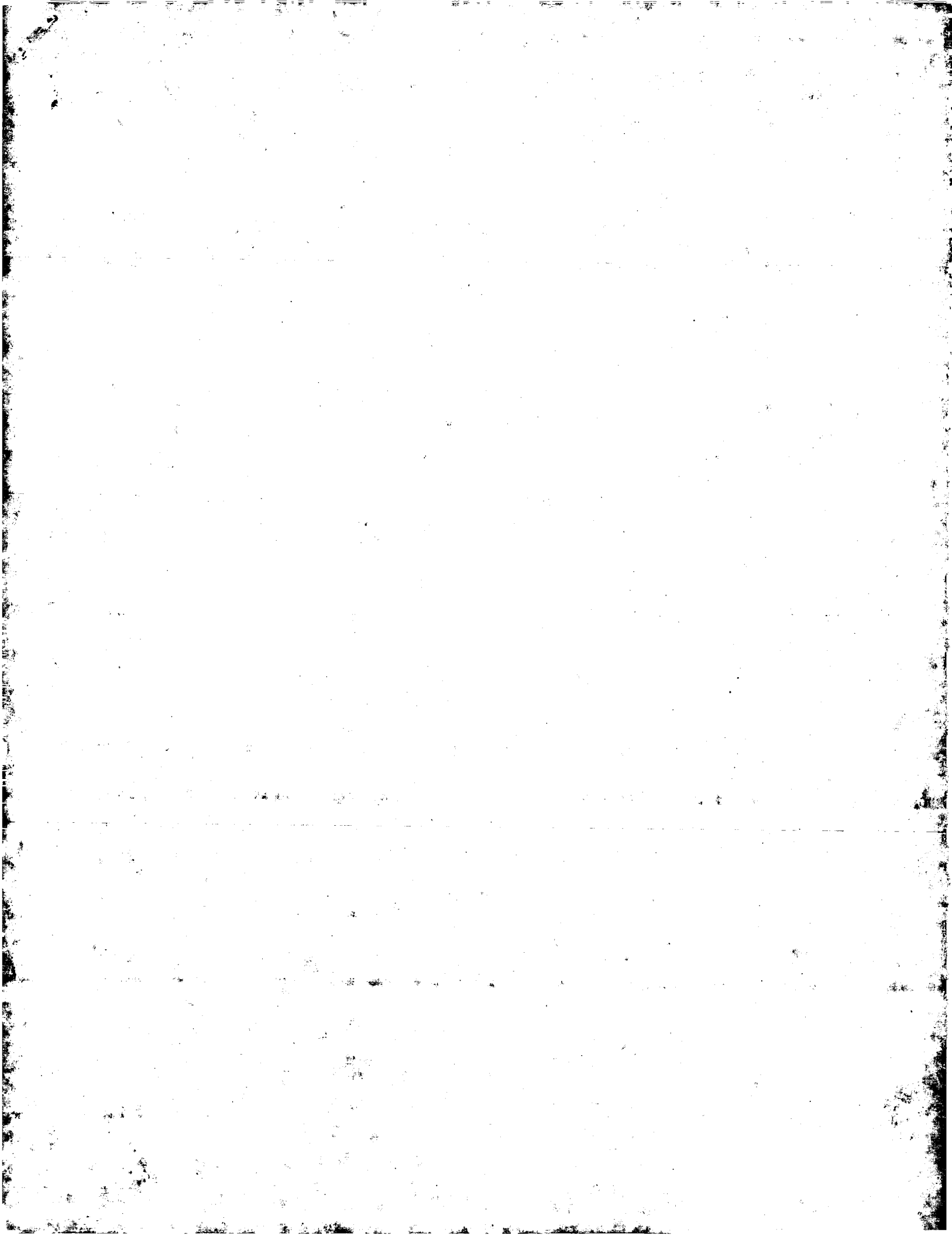
Query Match 30.7%; Score 159.5; DB 1; Length 218;
Best Local Similarity 47.7%; Pred. No. 0.00023;
Matches 52; Conservative 11; Mismatches 41; Indels 5; Gaps 3

QY 1 AKKYAKKAEKAYAKTAKAKKEKKAYAKK--EAKYNAAEAKKKAKAEAKKYAKKEAAKAK 57
   :||: :||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 109 SKKPGGLEKAPKKKLSAAKPKKAALKPPAALAKKPKKAVAVKKSPPKAKKPPAASATKS 168
   :||: :||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

QY 58 KEAYKAELK-KYAKAAKAEKKEYAA-AEAKAEAAKAYKAEAAKAAKE 104
   :|: ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 169 VKSPKKAAPKKAVAAKSPAKAKAVKPKAAKPKAAKPKAAKAKKAAK 217
   :||: :||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Search completed: January 21, 2004, 09:00:59
Job time : 15.8636 secs

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ID	O9MWX1	PRELIMINARY;	PRT; 372 AA.
DT	O9MWX1:		
DT	01-NOV-1999 (TREMBLrel. 12, Created)		
DT	01-NOV-1999 (TREMBLrel. 12, Last sequence update)		
DT	01-MAR-2003 (TREMBLrel. 23, Last annotation update)		
DE	TolA protein.		
DE	TOL.		
OS	Pseudomonas putida.		
OC	Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;		
OC	Pseudomonadaceae; Pseudomonas.		
OX	NCBI_TaxID=303;		
RN	[1]		
RP	SEQUENCE FROM N.A.		
RC	STRAIN-mt-2;		
RC	MEDLINE=96198174; PubMed=8626299;		
RA	Rodriguez-Herva J.J.; Ramos-Gonzalez M.I.; Ramos J.;		
RT	"The Pseudomonas putida peptidoglycan-associated outer membrane		
RT	lipoprotein (PAL) is involved in maintenance of the integrity of the		
RT	cell envelope."		
RL	J. Bacteriol. 178:1699-1706(1996).		
RN	[2]		
RP	SEQUENCE FROM N.A.		
RC	STRAIN-mt-2;		
RC	Ramos-Gonzalez I.;		
RL	Submitted (JUN-1995) to the EMBL/GenBank/DBJ databases.		
RP	SEQUENCE FROM N.A.		
RC	STRAIN-mt-2;		
RA	Rodriguez-Herva J.J.;		
RL	Submitted (JUL-1999) to the EMBL/GenBank/DBJ databases.		
RN	[4]		
RP	SEQUENCE FROM N.A.		
RC	STRAIN-mt-2;		
RC	MEDLINE=96422022; PubMed=8824639;		
RA	Rodriguez-Herva J.J.; Ramos J.;		
RT	"Characterization of an OptI null mutant of Pseudomonas putida.";		
RL	J. Bacteriol. 178:5836-5840(1996).		
DR	EMBL; X74218; CAB50780.1; -		
DR	InterPro; IPRO05819; Histone_H5.		
DR	InterPro; IPRO06260; TonB_C.		
DR	PRINTS; PRO0624; HISTONEH5.		
DR	TIGRFAMs; TIGR01352; tonB_Cterm; 1.		
SQ	SEQUENCE 372 AA; 4013 MW; 87F49785ECC3C0BC CRC64;		
Query Match	36.7%; Score 190.5; DB 2; Length 372;		
Best Local Similarity	50.4%; Pred. No. 6.3e-06;		
Matches	59; Conservative 14; Mismatches 31; Indels 13; Gaps 4		
OY	5 AKKAERAYAKKAAK---EKAYAKKEAKYYKAE-----AKKKAKAEKKYAKEAA 54		
Db	118 AKKADA-AKAAEAARKAAEAARKADEAKKAEKQADIAKKADEAKKAEBA 176		
OY	55 K--AKKEAYKAEKKYAYAKKEKKEEVAAAEKKAEEAKAYAAEAKAAKAEAYEA 109		
Db	177 KKAABEAKKKALEDKKAABEAKKKAEDAKKGAABEDAKKKAABEAKKKAADA 233		
RESULT 3			
OBTSC8	PRELIMINARY;	PRT; 1866 AA.	
AC	OBTSC8:		
DT	01-JUN-2002 (TREMBlrel. 21, Created)		
DT	01-JUN-2002 (TREMBlrel. 21, Last sequence update)		
DT	01-OCT-2002 (TREMBlrel. 22, Last annotation update)		
DE	MaebI.		
GN	MAEBI.		
OS	Plasmodium vivax.		
OC	Eukaryota; Alveolata; Apicomplexa; Haemosporidia; Plasmodium.		
OX	NCBI_TaxID=5885;		

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RN  [1]
RP  SEQUENCE FROM N.A.
RC  STRAIN=Salvador;
RA  Michon P., Stevens J.R., Kaneo O., Adams J.H.;
RT  "Evolutionary relationships of conserved cysteine-rich motifs in
RL  adhesive molecules of malaria parasites.";
DR  Mol. Biol. Evol. 0:0-0(2002).
SQ  EMBL; AY042083; AAL10508.1; -
    SEQUENCE 1866 AA; 212420 MW; DC692D7CFAE7D93F CRC64;

Query Match          36.7%; Score 190.5; DB 5; Length 1866;
Best Local Similarity 49.6%; Pred. No. 2,7e-05;
Matches 61; Conservative 17; Mismatches 22; Indels 23; Gaps 6;

QY  5 AKKAEKAYAKKAKAAKAE-----KKAAYKKEAKAYKAAEAKKKA-----KAEAKKKA 50
DB  1262 AKKAE--ARRAEAKKAEAEAKKAEAKKAAEAAKAEAAKKAEEARAEAEAKKAEAAKKA 1319
QY  51 KEAAKAKKEAAKAAKKYAKAKAAKKEKKEVAAAEAKKAEAAK-----AVKAEAAKAAAEAA 106
DB  1320 EDARKA--EARKAAEAAKKAEEARAEARAEAKKAEAAKKAEEAAKKAEEARAEAAKKA--EAA 1376
QY  107 YEAA 109
DB  1377 RKA 1379

RESULT 4
Q8FJT1 PRELIMINARY; PRT; 421 AA.
ID Q8FJT1
AC Q8FJT1
DT 01-MAR-2003 (TREMBLrel. 23, Created)
DT 01-MAR-2003 (TREMBLrel. 23, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE ToLA protein.
GN TOLA OR C0818.
OS Escherichia coli O6.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Escherichia.
OX NCBI_TaxID=217992;
RN  [1]
RP  SEQUENCE FROM N.A.
RC  STRAIN=O6:HI / CFT073 / ATCC 700928;
EX MEDLINE=22388234; PubMed=12471157;
RA Welch R.A., Burland V., Plunkett G. III, Redford P., Roesch P.,
RA Raeko D., Buckles E.J., Liou S.-R., Boutin A., Hackett U., Stroud D.,
RA Mayhew G.F., Rose D.J., Zhou S., Schwartz D.C., Perna N.T.,
RA Mobley H.L.T., Dommberg M.S., Blattner F.R.;
RT "Extensive mosaic structure revealed by the complete genome sequence
RT of uropathogenic Escherichia coli.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:17020-17024(2002).
DR EMBL; AE016757; AAN79291.1; -.
KW Complete proteome.
SQ  SEQUENCE 421 AA; 43184 MW; DB296626F056D385 CRC64;

Query Match          36.3%; Score 188.5; DB 16; Length 421;
Best Local Similarity 53.0%; Pred. No. 9.6e-06;
Matches 61; Conservative 11; Mismatches 32; Indels 11; Gaps 5;

QY  1 AKKAYKAAE-----FAVAKKAAAEKKAAYKAEAKKAAKKAEEAKKAAEAKKAAEAKK 55
DB  120 ABEAAKQAEELKQKQAEAEAAAGAAADAKKAADAKA--AEAAKKAAGAAADAKKAEAEAAK 177
QY  56 AKKEAY-KAEAKKYAKAKAEKKEVAAAEAAK-----AEAAKAYKAAEAAKAAEAA 106
DB  178 AAVEAAKKAEEAAALAKKKKAAEAAEAAAEAAAEAAKKAATEAAEAKKAEAEKAAAEKAA 232

RESULT 5
Q8ZOT6 PRELIMINARY; PRT; 407 AA.
ID Q8ZOT6
AC Q8ZOT6
DT 01-MAR-2002 (TREMBLrel. 20, Created)

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DR DNA Res 8:11-22 (2001).
DR EMBL: A6005252; AAG55075.1; -.
DR EMBL: AP002553; BAB34197.1; -.
DR InterPro: IPR000104, Antifreeze_1.
DR PRINTS: PR00308; ANTIFREEZE1.
KW Complete proteome.
SQ SEQUENCE 394 AA; 40517 MW; 5B58D8E8230BDE28 CRC64;

Query Match 35.0%; Score 181.5; DB 16; Length 394;
Best Local Similarity 52.2%; Pred. No. 2.7e-05;
Matches 60; Conservative 12; Mismatches 32; Indels 11; Gaps 5;

QY 1 AKKAKKAE---KAYAKKAAAEKKAYAKKEKAYAKAAEAKKKKATAEAKVKYK-ENAK 55
Db 120 AEEAAKQAEIKQKQAEAAAKAAADADAKQAEADDKA--AEAAKQAAADAKKQAEADAK 177
QY 56 AKKEAY-KAEAKKYAKAAKAEKKEKYAAAEAKK--AEAAAYKAAKAAKAAKAA 106
Db 178 AAEEAQKAEAAEAALAKKQAEAAEAARKKAAAEKAAADKKAEEKAAAEKAA 232

RESULT 7
Q828C1 PRELIMINARY; PRT; 376 AA.
AC Q828C1
DT 01-MAR-2002 (TrEMBLrel. 20, Created)
DT 01-MAR-2002 (TrEMBLrel. 20, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Tola Protein.
GN STY0793.
OS Salmonella typhi.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Salmonella.
OX NCBI_TaxID=601;
RX [1]
RP SEQUENCE FROM N.A.
RC STEAIN-CT18;
RA MEDLINE=21534947; PubMed=11677608;
RA Parkhill J., Dougan G., James K.D., Thomson N.R., Pickard D., Wain J.,
RA Churcher C., Mungall K.L., Bentley S.D., Holden M.T.G., Sebahia M.,
RA Baker S., Baaham D., Brooks K., Chillingworth T., Conerton P.,
RA Cronin A., Davis P., Davies R.M., Dowd L., White N., Farrar J.,
RA Feltham T., Hamlin N., Haque A., Hien T.T., Holtroyd S., Jaseles K.,
RA Krogh A., Larsen T.S., Leather S., Moule S., O'Gaora P., Parry C.,
RA Quail M., Rutherford K., Simmonds M., Skelton J., Stevens K.,
RA Whitehead S., Barrett B.G.;
RT "Complete genome sequence of a multiple drug resistant Salmonella
RT enterica serovar Typhi CT18."
RL Nature 413:848-852(2001).
RV EMBL: AL627268; CAD05209.1; -.
KW Complete proteome.
SQ SEQUENCE 376 AA; 38804 MW; EC21F2C4767A8A42 CRC64;

Query Match 34.3%; Score 178; DB 16; Length 376;
Best Local Similarity 52.3%; Pred. No. 4.5e-05;
Matches 58; Conservative 10; Mismatches 33; Indels 10; Gaps 4;

QY 1 AKKAKKAEAYAKKAAAEKKAYAKKEKAYAKAAEAKKKKATAEAKVKYKAEAAKKEA 60
Db 137 AKAAADAKKQAEAAKAAADADAKKQAEAEAKAAAEAKKAAAEAE--AKAAAEAKK-- 190
QY 61 YKAEAKYAKAAKAEKKEKYAAAEKKAEEAAAYKAAKAAKAA--KEAAVE 108
Db 191 -KAEABAAKAAADADAKKQAEAAKAAAEAKKQADAAKAAADAKKQAAAE 240

RESULT 8
Q82GZ2 PRELIMINARY; PRT; 388 AA.
AC Q82GZ2
DT 01-MAR-2002 (TrEMBLrel. 20, Created)
DT 01-MAR-2002 (TrEMBLrel. 20, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)

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DE TOLA colicin import membrane protein.
GN TOLA OR YP01123.
OS Yersinia pestis.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Yersinia.
NCBI_Taxid=632;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CO-92 / Biovar Orientalis;
RX MEDLINE=21470413; PubMed=11586360;
RA Parhill J., Wren B.W., Thomson N.R., Tibball R.W., Holden M.T.G.,
RA Prentice M.B., Sebahia M., James K.D., Churcher C., Mungall K.L.,
RA Baker S., Basham D., Bentley S.D., Brooks K., Cerdeno-Tarraga A.M.,
RA Chillingworth T., Cronin A., Davies R.M., Davis P., Dougan G.,
RA Felwell T., Hamlin N., Holroyd S., Jagsels K., Karlyshev A.V.,
RA Leather S., Moule S., Oyston P.C.F., Quail M., Rutherford A.,
RA Simmonds M., Skelton J., Stevens K., Whitehead S., Barrell B.G.;
RA "Genome sequence of Yersinia pestis, the causative agent of plague.",
RL Nature 413:523-527(2001).
DR EMBL; AJ414146; CAC89966.1; -.
KW Complete proteome.
SQ SEQUENCE 388 AA; 40424 MW; 81447B04B30A7E7C CRC64;

Query Match 34.3%; Score 178; DB 16; Length 388;
Best Local Similarity 52.1%; Pred. No. 4.6e-05;
Matches 63; Conservative 12; Mismatches 28; Indels 18; Gaps 7;

QY 2 KYAKKAEKAYKAKKAEK-----AYAKKEA-KAYKA-AEAKKAEKAYKAKA 53
DB 140 KQAEQOKIAAAAVAKAKKEQOKAETAAQAKAEADKIYKQAEQOKAEAKKAAVA 199
QY 54 AKAKKAYKAEKAYKAKKAEK-----AEKKEVAAAEKKAKEAK-AKYAEAKKAAKAA 105
DB 200 AAKKQA-DADAKKAVEAEKKAADAEKKAADAE-KKAAAKKVAATAAEKKAATAA 257
QY 106 A 106
DB 258 A 258

RESULT 9
Q8CZ28 PRELIMINARY; PRT; 393 AA.
AC Q8CZ28;
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Membrane spanning protein.
GN TOLA OR Y1056.
OS Yersinia pestis.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Yersinia.
NCBI_Taxid=632;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=KIMS / Biovar Mediaevalis;
RX MEDLINE=22137863; PubMed=21242430;
RA Deng W., Burland V., Plunkett G. III, Boutin A., Mayhew G.F., Lies P.,
RA Perna N.T., Rose D.J., Mau B., Zhou S., Schwartz D.C.,
RA Fetherston J.D., Lindler L.B., Brubaker R.R., Plano G.V.,
RA Straley S.C., McDonough K.A., Nilles M.L., Watson J.S., Blattner F.R.,
RA Perry R.D.;
RA "Genome sequence of Yersinia pestis KIM.",
RL J. Bacteriol. 184:4601-4611(2002).
DR EMBL; AB013906; AAM86606.1; -.
SQ SEQUENCE 393 AA; 41012 MW; 1E3E4F87B53481 CRC64;

Query Match 34.3%; Score 178; DB 16; Length 393;
Best Local Similarity 52.1%; Pred. No. 4.7e-05;
Matches 63; Conservative 12; Mismatches 28; Indels 18; Gaps 7;

QY 2 KYAKKAEKAYKAKKAEK-----AYAKKEA-KAYKA-AEAKKAEKAYKAKA 53

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DB 145 KQAEQOKIAAAAVAKAKKEQOKAETAAQAKAEADKIYKQAEQOKAEAKKAAVA 204
QY 54 AKAKKAYKAEKAYKAKKAEK-----AEKKEVAAAEKKAKEAK-AKYAEAKKAAKAA 105
DB 205 AAKKQA-DADAKKAVEAEKKAADAEKKAADAE-KKAAAKKVAATAAEKKAATAA 262
QY 106 A 106
DB 263 A 263

RESULT 10
Q8XV7 PRELIMINARY; PRT; 200 AA.
AC Q8XV7;
DT 01-MAR-2002 (TrEMBLrel. 20, Created)
DT 01-MAR-2002 (TrEMBLrel. 20, Last sequence update)
DT 01-MAR-2002 (TrEMBLrel. 23, Last annotation update)
DE Probable histone H1 protein.
DB RSC2793 OR R500453.
OS Ralstonia solanacearum (Pseudomonas solanacearum).
OC Bacteria; Proteobacteria; Betaproteobacteria; Burkholderiales;
OC Ralstoniaceae; Ralstonia.
NCBI_Taxid=305;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=GM11000;
RX MEDLINE=21681879; PubMed=11823852;
RA Salanoubat M., Genin S., Artiguenave F., Gouzy J., Mangenot S.,
RA Arlat M., Billault A., Broctier P., Gams J.C., Catolico L.,
RA Chandler M., Choisme N., Claudel-Renard C., Cunnac S., Demange N.,
RA Gaspier C., Lavie M., Moisan A., Robert C., Saurin W., Schlex T.,
RA Sigler P., Thebaud P., Whalen M., Wincker P., Levy M.,
RA Weisenbach J., Boucher C.A.;
RA "Genome sequence of the plant pathogen Ralstonia solanacearum.",
RL Nature 415:497-502(2002).
DR EMBL; AL646071; CAD16500.1; -.
DR InterPro; IPR005819; Histone_H5.
DR PRINTS; PR00624; HISTONEH5.
KW Complete proteome.
SQ SEQUENCE 200 AA; 19279 MW; D3831B590510272D CRC64;

Query Match 34.0%; Score 176.5; DB 16; Length 200;
Best Local Similarity 54.5%; Pred. No. 3.2e-05;
Matches 67; Conservative 5; Mismatches 32; Indels 19; Gaps 8;

QY 1 AKKVAKKAEKAYKAKKAEK-----KKAYAKKEAKAYKAAEK---KKA-----KAEKTY- 49
DB 42 AKKVA--AKKVAAKQAAPAKKAAVGVAAKKGAAPAKKAAVKKVAAKAPAKKAAVKKVA 99
QY 50 AKEAAKAKKAEKAYKAEKAYKAAK--AEKKEVAAAEKKAKEAKAYKAE--AKKAAKAAKAA 106
DB 100 AKKAPAKKAAVAKKVAKKAAKAPAKKAAK--APAKKAPAKKAAKAAKAPAKKAPAA 156
QY 107 YEA 109
DB 157 KKA 159

RESULT 11
Q9CM70 PRELIMINARY; PRT; 389 AA.
AC Q9CM70;
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE TOLA OR PM0968.
GN Pasteurella multocida.
OS Bacteria; Proteobacteria; Gammaproteobacteria; Pasteurellales;
OC Pasteurellaceae; Pasteurella.
NCBI_Taxid=747;
RN [1]

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RP SEQUENCE FROM N.A.
RC STRAIN=PM70;
RA MEDLINE=21145866; PubMed=11248100;
RA May B.J., Zhang Q., Li L.L., Paucetian M.L., Whittam T.S., Kapur V.;
RT "Complete genomic sequence of Pasteurella multocida PM70.";
RT Proc. Natl. Acad. Sci. U.S.A. 98:3460-3465(2001).
DR EMBL; AB06136; AAK03052.1; -.
DR HSSP; P19934; 1TOL.
DR InterPro; IPR000533; Tropomyosin.
DR PRINTS; PR00194; TROPOMYOSIN.
DR Complete proteome.
SQ SEQUENCE 389 AA; 42197 MW; B4032F2A2FD9E94B CRC64;

Query Match 33.4%; Score 173.5; DB 16; Length 389;
Best Local Similarity 48.7%; Pred. No. 9.4e-05;
Matches 55; Conservative 18; Mismatches 31; Indels 9; Gaps 4;

Qy 6 KKAERKAAKK-AKAAEK-KAYAKKEKAAKAAKKA--EAKKYAKA-----AKA 56
Db 145 KQAEKAKKQLAEMAKLKAEMAKRLAALAKQAEBAKKAEBEAKKAKAEKAKAEKAKA 204
Qy 57 KKEAYKAEKKYAKAKAEKKEVAAAEAKKAAKAYKAAKAAKAAKAAKAAKAA 109
Db 205 KVEKAKAEKAEKAKAEKAKAEKAKAEKAKAEKAKAEKAKAEKAKAEKAKA 257

RESULT 12
061164 PRELIMINARY; PRT; 1701 AA.
AC 061164;
DT 01-AUG-1998 (TREMBLrel. 07, Created)
DT 01-AUG-1998 (TREMBLrel. 07, Last sequence update)
DT 01-OCT-2002 (TREMBLrel. 22, Last annotation update)
DE Erythrocyte binding protein.
GN MAEBL.
OS Plasmodium yoelii yoelii.
OC Eukaryota; Alveolata; Apicomplexa; Haemosporida; Plasmodium.
OK NCBI_Taxid=73239;
RN 11
RP SEQUENCE FROM N.A.
RC STRAIN=YM;
RX MEDLINE=98115903; PubMed=9448314;
RA Kappe S.H.I., Noe A.R., Fraser T.S., Blair P.L., Adams J.H.;
RT "A family of chimeric erythrocyte binding proteins of malaria parasites.";
RT Proc. Natl. Acad. Sci. U.S.A. 95:1230-1235(1998).
RL DR EMBL; AF03186; AAC05366.1; -.
SQ SEQUENCE 1701 AA; 199268 MW; EDA8E2DEFD87C8BA CRC64;

Query Match 32.9%; Score 170.5; DB 5; Length 1701;
Best Local Similarity 53.5%; Pred. No. 0.00058;
Matches 54; Conservative 11; Mismatches 31; Indels 5; Gaps 5;

Qy 5 AKKAERKAAKKAAKAEKKAAYAKAEKKAAYAKAEKKAAYAKAEKKAAYAKA- 63
Db 1205 AKKAEEF-RKKAERAK-KAEAKKKAERAKKAEEERK-KAEAKKALEKKKSEAKKAL 1261
Qy 64 EAKKYAKAKAEKKEVAAAEAKKAAKAAKAAKAAKAAKAAKAAKAAKAAK 104
Db 1262 ERKKAERAKKAEKKAERAKKAAKAAKAAKAAKAAKAAKAAKAAKAAK 1301

RESULT 13
039576 PRELIMINARY; PRT; 232 AA.
AC 039576;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE Histone H1.
GN CH1.
OS Chlamydomonas reinhardtii.
OC Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;

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OC Chlamydomonadaceae; Chlamydomonas.
OK NCBI_Taxid=3055;
RN 11
RP SEQUENCE FROM N.A.
RX MEDLINE=96120862; PubMed=8590479;
RA Fabry S., Muller K., Lindauer A., Park P.B., Cornelius T., Schmitt R.;
RT "The organization structure and regulatory elements of Chlamydomonas histone genes reveal features linking plant and animal genes.";
RT Curr. Genet. 28:333-345(1995).
RL EMBL; U16726; AAA98452.1; -.
DR HSSP; P02259; 1HST.
DR InterPro; IPR005818; Histone H1/H5.
DR InterPro; IPR005819; Histone H5.
DR InterPro; IPR003216; Linkerhist N.
DR Pfam; PF00538; Linker_histone; I.
DR PRINTS; PR00624; HISTONEH5.
DR ProDom; PD000373; Linkerhist_N; 1.
DR SMART; SM00526; H15; 1.
SQ SEQUENCE 232 AA; 24693 MW; 2D006A44A8FA037 CRC64;

Query Match 32.8%; Score 170; DB 10; Length 232;
Best Local Similarity 46.3%; Pred. No. 0.0001;
Matches 57; Conservative 10; Mismatches 38; Indels 18; Gaps 5;

Qy 2 KKYAKKA---EKAYKKAKAKEKKAAYAKKEA---KAYKAAEAKKAAKAEKKAKEA- 53
Db 100 KKAARKAAATPKKAAAPKKEGAVKTKAPKKEGKPKAKAEKPKKEGKKAAKPAK 159
Qy 54 ---AKAKKEAYKAEKKYAKAKAEKKEVAAE---AKKAEPAK---AYKAAKAAKAAK 103
Db 160 AEKKRKAAPKAKPTTKTAAKAAKPKAEKKPKAAAKPKAEKPPKAAKPKAEKKAAPKAAK 219
Qy 104 EAA 106
Db 220 KSA 222

RESULT 14
001395 PRELIMINARY; PRT; 275 AA.
AC 001395;
DT 01-JUL-1997 (TREMBLrel. 04, Created)
DT 01-JUL-1997 (TREMBLrel. 04, Last sequence update)
DT 01-OCT-2002 (TREMBLrel. 22, Last annotation update)
DE Axoneme-associated protein MST101(3).
GN MST101(3) OR DHMST101.
OS Drosophila hydei (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Insecta; Pterygota;
OC Neoptera; Diptera; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
OK NCBI_Taxid=7224;
RN 11
RP SEQUENCE FROM N.A.
RA Neesen J., Heinlein U.A.O., Buenemann H.;
RL Submitted (JAN-1997) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: POSSIBLE STRUCTURAL ROLE IN THE SPERM TAIL (BY SIMILARITY).
CC -1- SUBCELLULAR LOCATION: CYTOPLASMIC (BY SIMILARITY).
CC -1- TISSUE SPECIFICITY: TESTIS (BY SIMILARITY).
CC -1- DOMAIN: THE PREDOMINANT STRUCTURE IS ALPHA-HELICAL.
DR EMBL; U85627; AAB51369.1; -.
DR FlyBase; FBgn020732; Dhyd1mar101(3).
DR KMW Repeat; Multigene family.
FT DOMAIN 64
FT 255
SQ SEQUENCE 275 AA; 30436 MW; 76BAA7B2A2DF732C CRC64;

Query Match 32.7%; Score 169.5; DB 5; Length 275;
Best Local Similarity 48.2%; Pred. No. 0.00013;
Matches 54; Conservative 16; Mismatches 29; Indels 13; Gaps 6;

Qy 1 AKYAKKAEKAYAKK--AKAAKKEKAYAKKAYKAAEAKKKAK-AEAKKYAKAEKAAK 57

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Db      51 AEDVKKCCBEAAAKKCAEAACEKEAAEK-----KCAEAACEKEAAEKKCAEAACEKE 106
Qy      58 KEAYKAEAKKVAKAARAEKKEVAAAARAK-AEAAKAYKAAAKAAKAAEAAE 108
Db      107 QEA--AQKKCCALAKEKE--AAKKCAEAACEKEAAEKKEAAEKCEAAAFK 153

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RESULT 15

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039598      PRELIMINARY;      PRT;      265 AA.
AC      039598;
DT      01-NOV-1996 (TrEMBLrel. 01, Created)
DT      01-NOV-1996 (TrEMBLrel. 01, last sequence update)
DT      01-OCT-2002 (TrEMBLrel. 22, last annotation update)
DE      Cgcr-4 product (Fragment).
GN      Cgcr-4.
OS      Chlamydomonas reinhardtii.
OC      Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
OC      Chlamydomonadaceae; Chlamydomonas.
OX      NCBI_Taxid=3055;
RN      [1]
RP      SEQUENCE FROM N.A.
RX      MEDLINE=92119224; PubMed=1731966;
RA      Wakarchuk W.W., Muller F.W., Beck C.F.;
RT      "Two GC-rich DNA elements of Chlamydomonas reinhardtii with complex
RT      arrangements of directly repeated sequence motifs."
RL      Plant Mol. Biol. 18:143-146(1992).
DR      EMBL; X17208; CAA35080.1; -.
FT      NON_TER
SQ      SEQUENCE      265 AA; 26216 MW; B5318B7377CF782 CRC64;

```

Query Match 31.6%; Score 164; DB 10; Length 265;
 Best Local Similarity 39.4%; Pred. No. 0.00029;
 Matches 50; Conservative 21; Mismatches 38; Indels 18; Gaps 2;

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Qy      1 AKKYAKKAEEA-----YAKKAKAAKKEKAYAKKAAKAAKAAKAAKAA 46
      79 AEAKEADEAEARAAEAARAAVAAEWAAAEAEARAAEAARAAEAARAAEAARAA 138
Qy      47 KKYAKAEAKKKEAYKAEAKKVAKAAK---AEKKEVAAAARAEAKKAEAKAYKAAKAA 102
Db      139 RVAAEARAAAEARAAAEARAAAEARAAAEARAAAEARAAAEARAAAEARAA 198
Qy      103 KEAAVEA 109
Db      199 EAKAKEA 205

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Search completed: January 21, 2004, 09:00:27
 Job time : 44.8831 secs

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Peptide #8 used in

XX	(TEVA-) TEVA PHARM USA INC.
PI	Gad A, Lis D;
DR	WPI; 2000-317499/27.
XX	
PT	Copolymer 1 related polypeptides used as molecular weight markers for
PT	glutiramer acetate and for treatment and prevention of immune diseases
XX	
PS	Claim 10; Page 14; 72pp; English.
XX	
CC	AA82571 to AA82577 represent specifically claimed copolymer molecular
CC	weight TV-marker polypeptides from the present invention. The present
CC	invention describes polypeptides (I) for determining the molecular
CC	weight of a copolymer (CP), which has an identified molecular weight
CC	and an amino acid composition corresponding to the copolymer. The
CC	polypeptides of the invention are used as molecular weight markers for
CC	glutiramer acetate related tetrapolymers. The polypeptides may also be
CC	used for treating and preventing immune diseases in a mammal. Autoimmune
CC	diseases which may be treated include either cell-mediated or
CC	antibody-mediated diseases. Such diseases include arthritic conditions,
CC	demyelinating diseases and inflammatory conditions, e.g. multiple
CC	sclerosis, rheumatoid arthritis, osteoarthritis, autoimmune haemolytic
CC	anaemia, autoimmune oophoritis, autoimmune thyroiditis, autoimmune
CC	uveoretinitis, Crohn's disease, chronic immune thrombocytopenia
CC	purpura, colitis, contact sensitivity disease, diabetes mellitus, Graves
CC	disease, Guillain-Barre's syndrome, Hashimoto's disease, idiopathic
CC	myxoedema, myasthenia gravis, psoriasis, pemphigus vulgaris, or systemic
CC	lupus erythematosus. Mediated diseases which can be treated
CC	include host-versus-graft disease, graft-versus-host disease, and
CC	delayed-type hypersensitivity. The polypeptides of the invention have
CC	defined molecular weights and physical properties which are analogous to
CC	glutiramer acetate molecules, which makes them ideal for use as
CC	molecular weight markers.
CC	
SQ	Sequence 109 AA:
OY	Query Match 100.0%; Score 519; DB 21; Length 109;
	Best Local Similarity 100.0%; Pred. No. 1.5e-36;
	Matches 109; Conservative 0; Mismatches 0; Indels 0; Gaps 0
DB	1 AKKYAKKKKAYAKKKAKEKKAYAKKBEAKKYKAAAEKKKAKKAYAKKAAKXKA 60
	1 AKKYAKKKKAYAKKKAKEKKAYAKKBEAKKYKAAAEKKKAKKAYAKKAAKXKA 60
OY	61 YKAEKKTKAKAACEKEEYAAAAYKAAEKAAKAYKKAEEAAAKAAEAAYA 109
	61 YKAEKKTKAKAACEKEEYAAAAYKAAEKAAKAYKKAEEAAAKAAEAAYA 109
DB	61 YKAEKKTKAKAACEKEEYAAAAYKAAEKAAKAYKKAEEAAAKAAEAAYA 109
RESULT 2	
ID	AA82576 standard; peptide; 86 AA.
AC	AA82576;
XX	
DT	28-JUL-2000 (first entry)
DE	
XX	Copolymer molecular weight TV-marker amino acid sequence SEQ ID NO:6.
KM	Copolymer; molecular weight marker; TV-marker; immune disease;
KM	glutiramer acetate; autoimmune disease; antiarthritic; neuroprotective;
KM	osteoporotic; immunosuppressive; antithyroid; antiinflammatory;
KM	antidiabetic; thyromimetic; haemostatic; antipsoriatic; dermatological;
KM	antiangemic; immunosuppressive; demyelinating disease; osteoarthritis;
KM	inflammatory condition; multiple sclerosis; rheumatoid arthritis;
KM	Crohn's disease; chronic immune thrombocytopenia purpura; colitis;
KM	diabetes mellitus; Graves disease; Guillain-Barre's syndrome; psoriasis;
KM	Hashimoto's disease; idiopathic myxoedema; myasthenia gravis;
KM	pemphigus vulgaris; systemic lupus erythematosus.
OS	Unidentified.

PN		WO200018794-A1.	
XX			
PD	06-APR-2000.		
XX			
PF	24-SEP-1999;	99WO-US22402.	
XX			
PR	25-SEP-1998;	98US-0101693.	
PA	(YEDA) YEDA RES & DEV CO LTD.		
PA	(TEVA-) TEVA PHARM USA INC.		
P1	Gad A, Lis D;		
DR	WPI: 2000-317499/27.		
XX			
PT	Copolymer 1 related polypeptides used as molecular weight markers for		
XX	glatiramer acetate and for treatment and prevention of immune diseases		
PS	Claim 10; Page 14; 72pp; English.		
XX			
CC	AA182571 to AA182577 represent specifically claimed copolymer molecular		
CC	weight TV-marker polypeptides from the present invention. The present		
CC	invention describes polypeptides (I) for determining the molecular		
CC	weight of a copolymer (Cp), which has an identified molecular weight		
CC	and an amino acid composition corresponding to the copolymer. The		
CC	polypeptides of the invention are used as molecular weight markers for		
CC	glatiramer acetate related tetrapolymers. The polypeptides may also be		
CC	used for treating and preventing immune diseases in a mammal. Autoimmune		
CC	diseases which may be treated include either cell-mediated or		
CC	antibody-mediated diseases. Such diseases include arthritic conditions,		
CC	demyelinating diseases and inflammatory conditions, e.g. multiple		
CC	sclerosis, rheumatoid arthritis, osteoarthritis, autoimmune haemolytic		
CC	anaemia, autoimmune oophoritis, autoimmune thyroiditis, autoimmune		
CC	uveoretinitis, Crohn's disease, chronic immune thrombocytopenia		
CC	purpura, colitis, contact sensitivity disease, diabetes mellitus, Graves		
CC	disease, Guillain-Barre's syndrome, Hashimoto's disease, idiopathic		
CC	myxoedema, myasthenia gravis, psoriasis, pemphigus vulgaris, or systemic		
CC	lupus erythematosus. Mediated-mediated diseases which can be treated		
CC	include host-versus-graft disease, graft-versus-host disease, and		
CC	delayed-type hypersensitivity. The polypeptides of the invention have		
CC	defined molecular weights and physical properties which are analogous to		
CC	glatiramer acetate molecules, which makes them ideal for use as		
CC	molecular weight markers.		
SQ	Sequence	86 AA;	
Query Match	60.2%; Score 312.5; DB 21; Length 86;		
Best Local Similarity	72.1%; Pred. NO. 2.2e-19;		
Matches	80; Conservative 1; Mismatches 3; Indels 27; Gaps 5		
OY	1 AKTAKKAKAYAKKAKAKKKAYAKKEAQAAYRAAEKKKKAKKAEAKKYAKKAQKKA 60		
DB	1 AKKYAKK-EKAYAKKA-----EKAKKKAEAAAYRAAEKKKK----- 36		
OY	61 YKAEKKKAKAKAEKKKYEAAAEK-KAEA-KAYKAEAAKAAEAAYEA 109		
DB	37 -KAEEKTKAKAKAEKKYEAAAEKTKAEAKKAYKAAKAAKAAEAAIEA 86		
RESULT 3			
ID	AA182575 standard; peptide; 77 AA.		
AC	AA182575;		
DT	28-JUL-2000 (first entry)		
DE	Copolymer molecular weight TV-marker amino acid sequence SEQ ID NO.5.		
XX			
KM	Copolymer; molecular weight marker; TV-marker; immune disease;		
KM	glatiramer acetate; autoimmune disease; antiarthritic; neuroprotective;		
KM	osteoporetic; immunosuppressive; antithyroid; antiinflammatory;		
KM	antidiabetic; thyromimetic; hemostatic; antipsoriatic; dermatological;		

XX	anti-inflammatory; immunosuppressive; demyelinating disease; osteoarthritis;
XX	inflammatory condition; multiple sclerosis; rheumatoid arthritis;
KW	Crohn's disease; chronic immune thrombocytopenia purpura; colitis;
KW	diabetes mellitus; Graves disease; Guillain-Barre's syndrome; psoriasis;
KW	Hashimoto's disease; idiopathic myxoedema; myasthenia gravis;
KW	periphagus vulgaris; systemic lupus erythematosus.
XX	
OS	Unidentified.
PN	
XX	WO200018794-A1.
XX	
PD	06-APR-2000.
XX	
PF	24-SEP-1999; 99WO-US22402.
XX	
PR	25-SEP-1998; 98US-0101693.
XX	
PA	(YEDA) YEDA RES & DEV CO LTD.
PA	(TEVA-) TEVA PHARM USA INC.
XX	
PI	Gad A, Lis D;
XX	
DR	WPI; 2000-317499/27.
PT	
PT	Copolymer 1 related polypeptides used as molecular weight markers for
PT	glutramer acetate and for treatment and prevention of immune diseases
XX	
PS	Claim 10; Page 14; 72pp; English.

CC AA182571 to AA182577 represent specifically claimed copolymer molecular
CC weight TV-marker polypeptides from the present invention. The present
CC invention describes polypeptides (I) for determining the molecular
CC weight of a copolymer (CP), which has an identified molecular weight
CC and an amino acid composition corresponding to the copolymer. The
CC polypeptides of the invention are used as molecular weight markers for
CC glutaric acetate related tetrapolymers. The polypeptides may also be
CC used for treating and preventing immune diseases in a mammal. Autoimmune
CC diseases which may be treated include either cell-mediated or
CC antibody-mediated diseases. Such diseases include arthritic conditions,
CC demyelinating diseases and inflammatory conditions, e.g. multiple
CC sclerosis, rheumatoid arthritis, osteoarthritis, autoimmune haemolytic
CC anaemia, autoimmune oophoritis, autoimmune thyroiditis, autoimmune
CC uveoretinitis, Crohn's disease, chronic immune thrombocytopaenia
CC purpura, colitis, contact sensitivity disease, diabetes mellitus, Graves
CC disease, Guillain-Barre's syndrome, Hashimoto's disease, idiopathic
CC myxoedema, myasthenia gravis, psoriasis, pemphigus vulgaris, or systemic
CC lupus erythematosus. Mediated-mediated diseases which can be treated
CC include host-versus-graft disease, graft-versus-host disease, and
CC delayed-type hypersensitivity. The polypeptides of the invention have
CC defined molecular weights and physical properties which are analogous to
CC glutaric acetate molecules, which makes them ideal for use as
CC molecular weight markers.

Sequence 77 AA;

Query Match	55.7%;	Score 289;	DB 21;	Length 77;
Best Local Similarity	67.0%;	Pred. No. 1.8e-17;		
Matches 73; Conservative	1;	Mismatches 3;	Indels 32;	Gaps 4;

Qy	1	AAGVAAKKA	EKAYAKTA	KAKEKKAA	KYAAKBEA	KYAAEA	KKKA	KA	EAKKYAEAA	AAKXEA	60
Dd	1	AKKVAKK	-EKAVAKKA-	-EKAARKA	EAAKYAKAEAKKA-	-----	-----	-----	-----	-----	36
Qy	61	YKAEAKTAKA	KAKEKEYYAA	AEAKKA	EAAKYYKKA	EAAKAA	KA	EAAK	EAAYA	109	
Dd	37	KAENKTAKA	KAAKEKEYYAA	EAK-EEAK--	--YKBAAPAA	KA	EAAVEA	YE	77		

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RESULT 4
AAY82574
ID AAY82574 standard; peptide; 66 AA
XX
AC AAY82574;

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XX	
DT	28-JUL-2000 (first entry)

DE	Copolymer molecular weight	TV-marker amino acid sequence	SEQ ID NO:4.
1	10000	10000	10000
2	10000	10000	10000
3	10000	10000	10000
4	10000	10000	10000
5	10000	10000	10000
6	10000	10000	10000
7	10000	10000	10000
8	10000	10000	10000
9	10000	10000	10000
10	10000	10000	10000
11	10000	10000	10000
12	10000	10000	10000
13	10000	10000	10000
14	10000	10000	10000
15	10000	10000	10000
16	10000	10000	10000
17	10000	10000	10000
18	10000	10000	10000
19	10000	10000	10000
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21	10000	10000	10000
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25	10000	10000	10000
26	10000	10000	10000
27	10000	10000	10000
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74	10000	10000	10000
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76	10000	10000	10000
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78	10000	10000	10000
79	10000	10000	10000
80	10000	10000	10000
81	10000	10000	10000
82	10000	10000	10000
83	10000	10000	10000
84	10000	10000	10000
85			

KX Copolymer; molecular weight marker; TV-marker; immune disease;
 KW osteoarthritis; autoimmune disease; antithyroid; neuroprotective;
 KW glycerol acetate; autoimmune disease; antithyroid; antinflammatory;
 KW glucocorticoid; immunosuppressive; antithyroid; antinflammatory;
 KW antidiabetic; thyromimetic; haemostatic; antipsychotic; dermatological;
 KW antidiabetic; immunosuppressive; demyelinating disease; osteoarthritis;
 KW inflammatory condition; multiple sclerosis; rheumatoid arthritis;
 KW Crohn's disease; chronic immune thrombocytopenia purpura; colitis;
 KW diabetes mellitus; Graves disease; Guillain-Barre's syndrome; psoriasis;
 KW Hashimoto's disease; idiopathic myxoedema; myasthenia gravis;
 KW pemphigus vulgaris; systemic lupus erythematosus

OS Unidentified.

PN WO200018794-A1

06-APR-2000
PD

PF 24-SEP-1999; 99WO-US22402.

PR 25-SEP-1998; 98US-0101693.

PA (YEDA) YEDA RES & DEV CO LTD.

XX 3

XX

XXI

XX Claim 10; Page 14; 72pp; English.

CC MAY82571 to MAY82577 represent

weight. V-model: polypeptides from the present invention. The present invention describes polypeptides (1) for determining the molecular weight of a copolymer (CP), which has an identified molecular weight and an amino acid composition corresponding to the copolymer. The polypeptides of the invention are used as molecular weight markers for glatimer acetate related retroviruses. The polypeptides may also be used for treating and preventing immune diseases in a mammal. Autoimmune diseases which may be treated include either cell-mediated or antibody-mediated diseases. Such diseases include arthritic conditions, demyelinating diseases and inflammatory conditions, e.g. multiple sclerosis, rheumatoid arthritis, osteoarthritis, autoimmune haemolytic anaemia, autoimmune oophoritis, autoimmune thyroiditis, autoimmune uveoretinitis, Crohn's disease, chronic immune thrombocytopaenia purpura, colitis, contact sensitivity disease, diabetes mellitus, Graves disease, Guillain-Barre's syndrome, Hashimoto's disease, idiopathic myxoedema, myasthenia gravis, psoriasis, pemphigus vulgaris, or systemic lupus erythematosus. Mediated-mediated diseases which can be treated include host-versus-graft disease, graft-versus-host disease, and delayed-type hypersensitivity. The polypeptides of the invention have defined molecular weights and physical properties which are analogous to glatimer acetate molecules, which makes them ideal for use as molecular weight markers.

Sequence 66 AA;

Query Match	44.0%	Score 228.5	DB 21	Length 66
Best Local Similarity	56.9%	Pred. NO. 1.8e-12		
Matches 62; Conservative	0	Mismatches 4	Indels 43	Gaps 4

QY 1 AKCTAKAEAYAYKKAAAEKKAYAKKEKAAYKAAAEAKKAAAEAKCTAKAEAAKKKEA 60
 ||||| ||||| ||||| |||||
 Db 1 AKCTAKK-EKAYAKAKKA-----EAKAKKA----- 25
 QY 61 YKAAKCTYAKAAKAEKKEYPAAAEKKAEAAAYKAEAPAKAAAEAYEA 109

CC The invention relates to a nucleic acid transport system (NTS) for
CC delivering nucleic acid into a cell. The NTS contains but is not limited
CC to 5 components: (a) the nucleic acid or a macromolecule to be delivered;
CC (b) a moiety that recognizes and binds to a cell surface receptor or
CC antigen or is capable of entering a cell through cytosis; (c) a nucleic
CC acid or macromolecular molecule binding moiety; (d) a moiety that is
CC capable of moving or initiating movement through a nuclear membrane; and
CC or (e) a lysins moiety that enables the transport of the entire complex
CC from the cell surface directly into the cytoplasm of the cell. The NTS
CC delivers nucleic acid into the cellular interior as well as the nucleus
CC of specific cells. The NTS can be used to treat disorders by targeting
CC specific nucleic acid accordingly. The NTS can also be used to create
CC transgenic animals for assessing human disease, such as cancer, in an

CC The sequence represents poly-Lys-Ala, used to bind nucleic acid in a
CC nucleic acid transporter system. The nucleic acid transporter system uses
CC nucleic acid binding complexes containing surface ligands which are
CC capable of binding to a cell surface receptor and entering the cell
CC through cytosol. The compounds of the invention are either ligands,
CC binding molecules (surface ligands), lysis agents, spacer molecules or
CC their intermediates. The ligands, binding molecules, lysis agents and
CC spacer molecules are used in nucleic acid transporter systems to deliver
CC nucleic acid into specific cells e.g. in gene therapy to deliver nucleic
CC acid into hepatocytes, muscle cells or bone forming cells.

[illegible]

XX	AA845852	standard; Protein; 100 AA.
XX	AA845852;	
DT	21-MAR-2001	(first entry)
DE	Nucleic acid transporter system peptide ligand SEQ ID NO 64.	
KW	Nucleic acid delivery; nucleic acid transporter system; hormone; enzyme;	
KW	growth factor; clotting factor; apolipoprotein; receptor; drug; oncogene;	
KW	tumor antigen; tumor suppressor; viral antigen; parasitic antigen;	
KW	bacterial antigen.	
OS	unidentified.	
PN	US6150168-A.	
XX	21-NOV-2000.	
PF	05-JUN-1995; 95US-0460971.	
PR	14-DEC-1993; 93US-0167641.	
PR	20-MAR-1992; 92US-0855389.	
PR	19-MAR-1993; 93WO-US02725.	
PA	(BAYU) BAYLOR COLLEGE MEDICINE.	
PI	Gottchalk S, Sparrow J, Cristiano RJ, Smith LC, Woo SLG;	
DR	WPI; 2001-049093/06.	
PT	Nucleic acid transporter system for delivering nucleic acid into a	
PT	cell, useful for delivering proteins and polypeptides to cells,	
PT	including growth factors, enzymes, hormones, and tumor suppressors -	
PS	Discloure; Column 125-126; 105pp; English.	
XX	This invention describes a novel system (I) for delivering a nucleic acid	
CC	to a cell, comprising a binding complex comprising a ligand binding	
CC	molecule noncovalently bound to a nucleic acid and covalently linked to a	
CC	surface ligand, and a second binding complex comprising a second binding	
CC	molecule noncovalently bound to a nucleic acid and covalently linked to a	
CC	nuclear ligand. The complexes are simultaneously bound to the nucleic	
CC	acid. The nucleic acid transporter system can also be used in a method	
CC	for the in vivo targeting of the insertion of DNA into a cell. It can	
CC	also be used in processes for producing transformed cell lines. The	
CC	system can be used to deliver a variety or proteins and polypeptides,	
CC	such as hormones, growth factors, enzymes, clotting factors,	
CC	apolipoproteins, receptors, drugs, oncogenes, tumor antigens, tumor	
CC	suppressors, viral antigens, parasitic antigens, and bacterial antigens.	
CC	The transporter system uses lysis agents to overcome the problems of	
CC	endosomal/lysosomal degradation seen with prior art systems.	
SO	Sequence 100 AA;	

Query Match 34.6%; Score 179.5; DB 22; Length 100;

Best Local Similarity 55.0%; Pred. No. 3.5e-08; Matches 55; Conservative 8; Mismatches 34; Indels 3; Gaps 3.

```

RESULT 12
ABJ18771 ID ABJ18771 standard; Protein; 347 AA.
AC ABJ18771;
AD
AE
AF
AG
AH
AI
AJ
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AM
AN
AO
AP
AQ
AR
AS
AT
AU
AV
AW
AX
AY
AZ
BA BB BC BD BE BF BG BH BI BJ BK BL BM BN BO BP BQ BR BS BT BU BV BW BX BY BZ CA CB CC CD CE CF CG CH CI CJ CK CL CM CN CO CP CQ CR CS CT CU CV CW CX CY CZ DA DB DC DD DE DF DG DH DI DJ DK DL DM DN DO DP DQ DR DS DT DU DV DW DX DY DZ EA EB EC ED EE EF EG EH EI EJ EK EL EM EN EO EP EQ ER ES ET EU EV EW EX EY EZ FA FB FC FD FE FF FG FH FI FJ FK FL FM FN FO FP FQ FR FS FT FU FV FW FX FY FZ GA GB GC GD GE GF GG GH GI GJ GK GL GM GN GO GP GQ GR GS GT GU GV GW GX GY GZ HA HB HC HD HE HF HG HH HI HJ HK HL HM HN HO HP HQ HR HS HT HU HV HW HX HY HZ IA IB IC ID IE IF IG IH II IJ IK IL IM IN IO IP IQ IR IS IT IU IV IW IX IY IZ JA JB JC JD JE JF JG JH JI JJ JK JL JM JN JO JP JQ JR JS JT JU JV JW JX JY JZ KA KB KC KD KE KF KG KH KI KJ KL KM KN KO KP KR KS KT KU KV KW KY KZ LA LB LC LD LE LF LG LH LI LJ LK LL LM LN LO LP LQ LR LS LT LU LV LW LX LY LZ MA MB MC MD ME MF MG MH MI MJ MK ML MN MO MP MQ MR MS MT MU MV MW MX MY MZ NA NB NC ND NE NF NG NH NI NJ NK NL NO NP NQ NR NS NT NU NV NW NX NY NZ OA OB OC OD OE OF OG OH OI OJ OK OL OM ON OP OQ OR OS OT OU OV OW OX OY OZ PA PB PC PD PE PF PG PH PI PJ PK PL PM PN PO PP PQ PR PS PT PU PV PW PX PY PZ QA QB QC QD QE QF QG QH QI QJ QK QL QM QN QO QQ QR QS QT QU QV QW QX QY QZ RA RB RC RD RE RF RG RH RI RJ RK RL RM RN RO RP RQ RR RS RT RU RV RW RX RY RZ SA SB SC SD SE SF SG SH SI SJ SK SL SM SN SO SP SQ SR SS ST SU SV SW SX SY SZ TA TB TC TD TE TF TG TH TI TJ TK TL TM TN TO TP TQ TR TS TT TU TV TW TX TY TZ UA UB UC UD UE UF UG UH UI UJ UK UL UM UN UO UP UQ UR US UT UU UV UW UX UY UZ VA VB VC VD VE VF VG VH VI VJ VK VL VM VN VO VP VQ VR VS VT VU VW VX VY VZ WA WB WC WD WE WF WG WH WI WJ WK WL WM WN WO WP WQ WR WS WT WU WV WW WX WY WZ XA XB XC XD XE XF XG XH XI XJ XK XL XM XN XO XP XQ XR XS XT XU XV XW XX XY XZ YA YB YC YD YE YF YG YH YI YJ YK YL YM YN YO YP YQ YR YS YT YU YV YW YX YY YZ ZA ZB ZC ZD ZE ZF ZG ZH ZI ZJ ZK ZL ZM ZN ZO ZP ZQ ZR ZS ZT ZU ZV ZW ZX ZY ZZ
ABJ18771 27-FEB-2003 (first entry)
DE Pseudomonas aeruginosa biofilm formation-related protein #35.
KM Biofilm formation modulation; biofilm-associated disease;
KW cystic fibrosis; AIDS; middle ear infection; acne; periodontal disease;
XX catheter-associated infection; medical device-associated infection.
OS Pseudomonas aeruginosa.
PN WO200265295-A2.
PD 31-OCT-2002.
PF 19-APR-2002; 2002WO-US12532.
PR 20-APR-2001; 2001US-285190P.
PR 24-OCT-2001; 2001US-344142P.
XX (IOWA ) UNIV IOWA RES FOUND.
PA (HARD ) HARVARD COLLEGE.
PI Whiteley M, Bangera MG, Lory S, Greenberg BP;
DR WPI; 2003-075601/07.
DN N-PsDB; ABT14593.
PT Identifying compound capable of modulating biofilm formation by
PR bacteria/bacterial antibiotic resistance, useful for treatment of
PT biofilm associated disease -
PS Claim 1; Page 119-120; 154pp; English.
CC The invention comprises a method for identifying a compound capable of
CC modulating biofilm formation by bacteria. The method of the invention is
CC useful for identifying a compound capable of modulating biofilm formation
CC by bacteria or modulating bacterial antibiotic resistance. The method of
CC the invention is also useful for diagnosing and treating a subject
CC (especially an immunocompromised human) that is afflicted with a biofilm-
CC associated disease or disorder, such as: cystic fibrosis; AIDS; middle
CC ear infections; acne; periodontal disease; catheter-associated
CC infections; and medical device-associated infections. The present amino
CC acid sequence represents a protein that is used in the invention.
SQ Sequence 347 AA;
Query Match 34.5%; Score 179; DB 24; Length 347;
Beet Local Similarity 49.1%; Pred. No. 1.4e-07;
Matches 54; Conservative 17; Mismatches 23; Indels 10; Gaps 4
QY 6 KKAERAAVKKAAKAEK-----RAYAKKEAAVYAAEAERKKAAAEKRYAKEAAKAKE 59
DB 99 QKLEGQQYAAAKAAEQKKADEARQAENQKAAAEKKADEAKYAEEKAAAEQKKQNDIAKR 158
QY 60 AYKAEAKKYVAQAARAEKEVEYAAAEKKAEEAAKAYAEAAKAAAEAAVEA 109

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Db 159 A-EDAKK--KAAEDAKK--AAEDAKKAAEAKKAAEAKKAAVEA 204

RESULT 13

ID AAY14928 standard; protein; 223 AA.

AAV14928;

25-OCT-1999 (first entry)

DE Amino acid sequence of M. vaccae antigen GV-45.

XX Mycobacterium vaccae protein; antigen; T cell activation; cytokine;
XX dendritic cell maturation; infectious disease; immune disorder; cancer;
XX respiratory system; mycobacterial infection; allergy; tuberculosis;
XX leprosy; sarcoidosis; lung cancer; asthma; skin disorder; psoriasis;
XX dermatitis; eczema; alopecia areata; skin cancer; basal carcinoma;
XX squamous cell carcinoma; melanoma.

OS Mycobacterium vaccae.

XX WO9932634-A2.

PD 01-JUL-1999.

PF 23-DEC-1998; 98WO-NZ00189.

XX 04-DEC-1998; 98US-0205426.

PR 23-DEC-1997; 97US-0996624.

PR 23-DEC-1997; 97US-0997080.

PR 11-JUN-1998; 98US-0095855.

PR 17-SEP-1998; 98US-0156181.

PA (GENE-) GENESIS RES & DEV CORP LTD.

XX Prestige RL, Skinner MA, Tan P, Visser ES, Watson J;

XX WPI; 1999-430163/36.

XX N-PSDB; AAT11393.

XX Enhancing immune response to an antigen

XX Claim 1; Page 239; 243p; English.

XX The invention provides heat-killed Mycobacterium vaccae, or recombinant
XX M. vaccae proteins. The M. vaccae proteins may be employed to activate
XX T cells and natural killer cells, to stimulate the production of
XX cytokines, to enhance the expression of co-stimulatory molecules on
XX dendritic cells and monocytes, and to enhance dendritic cell maturation
XX and function. The proteins can be expressed by standard recombinant
XX methodology. Pharmaceutical compositions comprising the proteins or
XX nucleic acid sequences encoding the proteins can be used for the
XX treatment, prevention, and detection of disorders including infectious
XX diseases, immune disorders and cancer. In particular, the compounds and
XX methods are used for treatment of diseases of the respiratory system,
XX such as mycobacterial infections, asthma, allergies, tuberculosis,
XX leprosy, sarcoidosis and lung cancer, and disorders of the skin such as
XX psoriasis, atopic dermatitis, eczema, allergic contact dermatitis,
XX alopecia areata, and skin cancers such as basal carcinoma, squamous cell
XX carcinoma and melanoma.

XX Sequence 223 AA;

Query Match 34.3%; Score 178; DB 20; Length 223;

Best Local Similarity 54.5%; Pred. No. 1.1e-07;

Matches 60; Conservative 6; Mismatches 34; Indels 10; Gaps 5;

1 AKKVKKK--AKKAVKKKAAKAKE---KKAVAKKKAAYKAAKAAKAAKAAKAAK 55
112 AKKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAK 170

Qy 56 AKKAVKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAK 103
Db 171 AKKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAK 218

RESULT 14

ID AAR84568 standard; Protein; 643 AA.

AAAR84568;

09-MAY-1996 (first entry)

XX Trypanosoma cruzi TCR27 polypeptide, Ag15.

XX TCR27; Chagas disease; repeat unit; diagnosis; blood screening;

XX recombinant; fusion protein; glutathione-S-transferase.

XX Trypanosoma cruzi.

XX Key Location/Qualifiers

XX Region 329..552

XX /label= repeat region

XX /note= "16 of 69 repeat units of 14 amino acids"

XX WO9525797-A1.

XX 28-SEP-1995.

XX 20-MAR-1995; 95WO-US03191.

XX 24-MAR-1994; 94US-0216894.

XX (KIRC/) KIRCHHOFF L V.

XX (OTSU/) OTSU K.

XX Kirchoff LV, Otsu K;

XX WPI; 1995-344618/44.

XX N-PSDB; AAT05332.

XX New polypeptide(s) from the TCR27 protein of Trypanosoma cruzi - as

XX immunosay reagent for specific diagnosis of Chagas disease, also

XX related nucleic acid and transformed cells

XX Disclosure; Page 40-41; 68pp; English.

XX AAR84565-R84569 are polypeptides of the TCR27 protein of T. cruzi
XX The proteins are all fusion products with glutathione-S-transferase
XX (GST) and some contain a linker sequence. The TCR27 protein comprises
XX a 95 amino acid (aa) N-terminal region; 69 repeats of a highly
XX conserved 14 aa C-terminal region; and a 68 aa C-terminal region. This sequence
XX encodes the GST sequence, the Ag44 polypeptide contg. 16 of the 69
XX repeat units and also contains the amino and carboxy terminal
XX peptides of TCR27. The TCR27 polypeptides of the invention are useful
XX for the diagnosis of Chagas disease (American Trypanosomiasis), they
XX are capable of detecting anti-T. cruzi antibodies; or for blood
XX screening. The TCR27 protein has epitopes to which most T. cruzi
XX infected individuals have antibodies. The TCR27 polypeptides will not
XX react with serum from patients with leishmaniasis, schistosomiasis,
XX or autoimmune disease and are hence less likely to cause false
XX positives in diagnosis.

XX Sequence 643 AA;

Query Match 31.3%; Score 162.5; DB 16; Length 643;

Best Local Similarity 44.4%; Pred. No. 6.5e-06;

Matches 52; Conservative 18; Mismatches 36; Indels 11; Gaps 4;

1 AKKVKKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAA 56
445 AAAR84568AAAR84568AAAR84568AAAR84568AAAR84568AAAR84568 504

Job time : 53.3766 secB

QY 57 KK--EAYKAEAKKYAKAKAEKKEVA-----AAEAKKAEAAKAYK-AAEAKKAAAEAA 106
 DB 505 TKVAEAEKQKAAEATKVAAEAEKQKAAEATKVAAEAEKQKAAEATKVAAEAEKQKAAEAA 561

RESULT 15

AAV34068
 ID AAV34068 standard; peptide; 158 AA.

AAV34068;

23-NOV-1999 (first entry)

XX Histone H1 isoform H1.5 PANCA-reactive fragment (residues 69-226).

XX Ulcerative colitis; histone; H1-like antigen; porin antigen; human;
 KW Bacteroides antigen; inflammatory bowel disease; IBD; PANCA; diagnosis;
 KM perinuclear anti-neutrophil cytoplasmic antibody; isoform.

XX Homo sapiens.

PN MO9945955-A1.

PD 16-SEP-1999.

XX 12-MAR-1999; 99WO-US05492.

XX 12-MAR-1998; 98US-0041889.

PA (REGC) UNIV CALIFORNIA.

PI Braun J, Cohavy O;

XX WPI; 1999-551215/46.

XX Use of histone H1, porin or Bacteroides antigens as targets for the
 PT diagnosis, prevention and treatment of ulcerative colitis -

PS Example 4; Page 125-126; 134pp; English.

CC The invention provides a method for the diagnosis, prevention and
 CC treatment of ulcerative colitis (UC) using histone H1-like antigen, a
 CC porin antigen or a Bacteroides antigen as a target antigen. The novel
 CC method of diagnosing UC in a subject suspected of having inflammatory
 CC bowel disease (IBD) comprises: (1) obtaining a sample from the subject;
 CC (2) contacting the sample with a histone H1-like antigen, or perinuclear
 CC anti-neutrophil cytoplasmic antibody (PANCA)-reactive fragment, to form a
 CC complex of the histone H1-like antigen, or the PANCA-reactive fragment,
 CC and antibody to the histone H1-like antigen; and (3) detecting the
 CC presence or absence of the complex; where the presence of the complex
 CC indicates that the subject has UC. The PANCA-reactive histone H1-like
 CC antigen, porin antigen and Bacteroides antigen are useful in the
 CC diagnosis, prevention and treatment of UC. The methods can also be used
 CC for identifying agents useful for treating UC. The present sequence
 CC represents a human histone H1 isoform H1.5 PANCA-reactive fragment.

XX Sequence 158 AA;

Query Match 30.9%; Score 160.5; DB 20; Length 158;
 Best Local Similarity 46.8%; Pred. No. 2.2e-06;
 Matches 51; Conservative 7; Mismatches 44; Indels 7; Gaps 2;

QY 1 AKKYAKKAEKAYAKKAKAEKKAAYAKKAEAAKAA-----EAKKKAAYAKKAEKAEAAK 55
 DB 51 AKPKAKKAGAAKAKKPAAGATPKK--AKKAAGAKKAVKTKPKKAKKPAAGVKKVAKSPK 108

QY 56 AKKEAYKAEAKKYAAKAAAEKKEKYAAAEAKKAEAAKAYKAEAAKAAAE 104
 DB 109 AKAAKPKKATKSPAKPKAVKPKAKPKAAKPKAAKPKAAKPKAAKAAAK 157

Search completed: January 21, 2004, 08:59:10

RESULT 2
 US-09-816-989A-6
 Sequence 6, Application US/09816989A
 Patent No. US20020115103A1
 GENERAL INFORMATION:
 APPLICANT: Gad, Alexander
 APPLICANT: Lis, Boris
 TITLE OF INVENTION: COPOLYMER 1 RELATED POLYPEPTIDES FOR USE AS MOLECULAR WEIGHT MARKERS
 TITLE OF INVENTION: AND FOR THERAPEUTIC USE
 FILE REFERENCE: 2609/60807-A-PCT-US
 CURRENT APPLICATION NUMBER: US/09/816, 989A
 CURRENT FILING DATE: 2001-03-23
 PRIOR APPLICATION NUMBER: 60/101, 693
 PRIOR FILING DATE: 1998-09-25
 PRIOR APPLICATION NUMBER: PCT/US99/22402
 PRIOR FILING DATE: 1999-09-24
 NUMBER OF SEQ ID NOS: 7
 SOFTWARE: PatentIn version 3.1
 SEQ ID NO 6
 LENGTH: 86
 TYPE: PRT
 ORGANISM: Artificial Sequence
 FEATURE:
 OTHER INFORMATION: Description of Artificial Sequence: Synthetic Peptide
 US-09-816-989A-6

Query Match	60.2%	Score 312.5;	DB 10;	length 86;
Best Local Similarity	72.1%;	Pred. No. 3.2e-18;		
Matches 80; Conservative	1;	Mismatches 3;	Indels 27;	Gaps 5;

Oy 1 AKRYAKKA EKAYAKKAAKEKKA YAKKEKAYYAAAEAKKAKA EAETKAYAEAKKAKEA 60
||| ||| ||| : ||| ||| |||
Dd 1 AKKYAKK-EKAYAKKA-----EKAKKA EAKYAKAEAKKKA----- 36

Qy 61 YKAARKYAKAAKAEKEYYAAAEAK -KAEAA-KAYTAFAAKAALAKEAAYEA 109
| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 37 -KAEAKKYAKAAKAEKEYYAAAEAKYKAEEAKKAYTAEAAKAAKAAKEAAYEA 86

RESULT 3
US-09-816-989A-5
; Sequence 5, Application US/09816989A
; Patent No. US20020115103A1

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: APPLICANT: lis, Doris
:
: TITLE OF INVENTION: COPOLYMER 1 RELATED POLYPEPTIDES FOR USE AS MOLECULAR WEIGHT MARK
:
: TITLE OF INVENTION: AND FOR THERAPEUTIC USE

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; PRIOR APPLICATION NUMBER: PCT/US99/22402
; PRIOR FILING DATE: 1999-09-24
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 5
; LENGTH: 77
; TYPE: PPT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic Peptide
; US-09-816-989A-5

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Query Match	55.7%;	Score 289;	DB 10;	Length 77;
Best Local Similarity	67.0%;	Pred. No. 2.1e-16;		
Matches	73;	Conservative	1;	Mismatches 3;
			Indels	32;
			Gaps	4

Qy 1 AKKYAKKA EKAYAKKAAKKEKAYAKKEKAYAKKAAEAKKAKA EAKKYAEAKAKKEA 60

Db 1 AKKYAKK-EKAYAKKA-----EKAKKAAKAYAKKAAEKKA----- 36

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QY      61 YKAEAKKYAKAAKAKEKKEYYAAAEAKKAEEAAKAYKAEAAKAAAKEAAYEA 109
        |||||
Db       37 -KAEAKKYAKAAKAKEKKEYYAAAEAK-----YKAEAAKAAAKEAAYEA 77
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RESULT 4
US-09-816-989A-4
Sequence 4 Amplification US/09816989A

Patent No. US20020115103A1
; GENERAL INFORMATION:

APPLICANT: Gad, Alexander
 APPLICANT: Lis, Doris
 TITLE OF INVENTION: COPOLYMER 1 RELATED POLYPEPTIDES FOR USE AS MOLECULAR WEIGHT MARKERS
 TITLE OF INVENTION: AND FOR THERAPEUTIC USE

FILE REFERENCE: 2609/60807-A-PCT-US
CURRENT APPLICATION NUMBER: US/09/816,989A
CURRENT FILING DATE: 2001-03-23
PRIOR APPLICATION NUMBER: 60/101,693
PCT FILING DATE: 2000-03-07

PRIOR FILING DATE: 1998-09-29
PRIOR APPLICATION NUMBER: PCT/US99/22402
PRIOR FILING DATE: 1999-09-24

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; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: PatentIn version 3.11
; SEQ ID NO 4

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; LENGTH: 66
; TYPE: PRT
;

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; ORGANISM: Artificial Sequence
;
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic Peptide
US-09-816-989A-4

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Query Match	44.0%	Score 238.5	DB 10	length 66
Best Local Similarity	56.9%	Pred. No. 1	1e-11	
Matches 62	Conservative 0	Mismatches 4	Indels 43	Gaps 4

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Qy      1 AKCVAKKAEEAYAKCKAAKEKAYAKKEEAKAYAAEAARAKKAAKAKVAKKEAKKEA 60
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db      1 AKCVAKK-EKAVAKAKKA-----EAKAKKA----- 25

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QY      61 YKAAKKYAKAKAKEKYEAAAEEKKAEAAKAYKAEAAKAAKAEAYEA 109
        |||||
Db       26 -KAAKKYAKAKAKEKYEAAAEEK-----YKAEAAKAAKAEAYEA 66
```

RESULT 5
US-09-816-989A-3

Patent No. US20020115103A1
; GENERAL INFORMATION:

APPLICANT: Gad, Alexander
APPLICANT: Lis, Doris
TITLE OF INVENTION: COPOLYMER 1 RELATED POLYPEPTIDES FOR USE AS MOLECULAR WEIGHT MARKERS
TITLE OF INVENTION: AND FOR THERAPEUTIC USE

FILE REFERENCE: 2609/60807-A-PCT-US
CURRENT APPLICATION NUMBER: US/09/816,989A
CURRENT FILING DATE: 2001-03-23
PRIOR APPLICATION NUMBER: 60/101,693

PRIOR FILING DATE: 1998-09-25
PRIOR APPLICATION NUMBER: PCT/US99/22402
PRIOR FILING DATE: 1999-09-24

```

; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: PatentIn version 3.1.1

```

```

; SEQ ID NO 3
;
; LENGTH: 56
;
; TYPE: PRT
;
; ORGANISM: Artificial Sequence
;
; PRATM:

```

OTHER INFORMATION: Description of Artificial Sequence: Synthetic Peptide

Query Match	34.8%;	Score 180.5;	DB 10;	Length 56;
Best Local Similarity	45.9%;	Pred. No. 5.8e-08;		

```

RESULT 7
US-10-205-979-52
Sequence 52, Application US/10205979
Publication No. US20030147861A1
GENERAL INFORMATION:
APPLICANT: Watson, James D.
APPLICANT: Tan, Paul L. J.
APPLICANT: Abernethy, Nevyn
TITLE OF INVENTION: Compounds and Methods for the Modulation
of Immune Responses
FILE REFERENCE: 11000.1063U
CURRENT APPLICATION NUMBER: US/10/205,979
CURRENT FILING DATE: 2002-07-25
PRIOR APPLICATION NUMBER: 60/308,446
PRIOR FILING DATE: 2001-07-26
NUMBER OF SEQ ID NOS: 52
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 52
LENGTH: 223
TYPE: prt
ORGANISM: Mycobacterium vaccae
US-10-205-979-52

```

RESULT 9
US-1056-761-9889
Sequence 9889, Application US/10156761
Publication No. US20030119018A1
GENERAL INFORMATION:
APPLICANT: OMURA, SATOSHI
APPLICANT: IKEDA, HARUO
APPLICANT: ISHIKAWA, JUN
APPLICANT: HORIKAWA, HIROSHI
APPLICANT: SHIBA, TADAYOSHI
APPLICANT: SAKAKI, YOSHIYUKI
APPLICANT: HATTORI, MASAHIRA
TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
FILE REFERENCE: 249-262
CURRENT APPLICATION NUMBER: US/10/156,761
PRIORITY FILING DATE: 2002-05-29
PRIORITY APPLICATION NUMBER: JP 2001-204089
PRIORITY FILING DATE: 2001-05-30
PRIORITY APPLICATION NUMBER: JP 2001-272697
PRIORITY FILING DATE: 2001-06-02
NUMBER OF SEQ ID NOS: 15109
SEQ ID NO 9889

LENGTH: 376
TYPE: PRT
ORGANISM: Streptomyces avermitilis
US-10-156-761-9889

Query Match 31.4%; Score 163; DB 15; Length 376;
Best Local Similarity 44.9%; Pred. No. 9.9e-06;
Matches 48; Conservative 15; Mismatches 38; Indels 6; Gaps 2;

QY 6 KKAAYAKKAAKAAK---KKAAYAKKAAKAAK---KKAAYAKKAAKAAK 59
DB 245 RRAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAK 304

QY 60 AKKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAK 106
DB 305 AKKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAK 351

RESULT 10
US-09-820-843A-27
Sequence 27, Application US/09820843A
Publication No. US2003003963A1
GENERAL INFORMATION:

APPLICANT: Council of Scientific and Industrial Research
TITLE OF INVENTION: A COMPUTATIONAL METHOD FOR THE IDENTIFICATION OF CANDIDATE PROTEI
FILE REFERENCE: Q63915
CURRENT APPLICATION NUMBER: US/09/820,843A
CURRENT FILING DATE: 2001-03-30
NUMBER OF SEQ ID NOS: 118
SOFTWARE: Patent version 3.0
SEQ ID NO 27
LENGTH: 356
TYPE: PRT
ORGANISM: Vibrio cholerae
FEATURE:
NAME/KEY: misc. feature
OTHER INFORMATION: colA protein
NAME/KEY: misc. feature
OTHER INFORMATION: gII9656364
US-09-820-843A-27

Query Match

Best Local Similarity 31.2%; Score 162; DB 11; Length 356;
Matches 55; Conservative 13; Mismatches 37; Indels 24; Gaps 5;

QY 3 KKAAYAKKAAKAAK---AKAAYAKKAAKAAKAAKAAK---AKAAYAKKAAKAAK 56
DB 112 KKAAYAKKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAK 171
QY 57 KKAAYAKKAAKAAK---AKAAYAKKAAKAAKAAKAAKAAKAAKAAK 100
DB 172 KKAAYAKKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAK 229
QY 101 AKKAAKAAKAAK 109
DB 230 AKKAAKAAKAAK 238

RESULT 11
US-10-229-567-40
Sequence 40, Application US/10229567
Publication No. US20030092080A1
GENERAL INFORMATION:

APPLICANT: Braun, Jonathan
TITLE OF INVENTION: Diagnosis, Prevention and Treatment of
Ulcerative Colitis, and Clinical Subtypes Thereof, Using
Microbial UC PANCA antigens
NUMBER OF SEQUENCES: 41
CORRESPONDENCE ADDRESS:
ADDRESSEE: Campbell & Flores LLP
STREET: 4370 La Jolla Village Drive, Suite 700

CITY: San Diego
STATE: California
COUNTRY: USA
ZIP: 92122

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/10/229,567
FILING DATE: 27-Aug-2002
CLASSIFICATION: <Unknown>

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/09/417,264
FILING DATE: <Unknown>
APPLICATION NUMBER: US 09/041,889
FILING DATE: <Unknown>

ATTORNEY/AGENT INFORMATION:
NAME: Campbell, Cathryn A.
REGISTRATION NUMBER: 31,815
REFERENCE/DOCKET NUMBER: P-PW 3006
TELECOMMUNICATION INFORMATION:
TELEPHONE: (619) 535-9001
TELEFAX: (619) 535-8949

INFORMATION FOR SEQ ID NO: 40:
SEQUENCE CHARACTERISTICS:
LENGTH: 158 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
SEQUENCE DESCRIPTION: SEQ ID NO: 40:
US-10-229-567-40

Query Match 30.9%; Score 160.5; DB 15; Length 158;
Best Local Similarity 46.8%; Pred. No. 6.4e-06;
Matches 51; Conservative 7; Mismatches 44; Indels 7; Gaps 2;

QY 1 AKKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAK 55
DB 51 AKKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAK 108

QY 56 AKKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAK 104
DB 109 AKKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAKAAK 157

RESULT 12
US-10-229-567-32

Sequence 32, Application US/10229567
Publication No. US20030092080A1
GENERAL INFORMATION:

APPLICANT: Braun, Jonathan
TITLE OF INVENTION: Diagnosis, Prevention and Treatment of
Ulcerative Colitis, and Clinical Subtypes Thereof, Using
Microbial UC PANCA antigens
NUMBER OF SEQUENCES: 41
CORRESPONDENCE ADDRESS:
ADDRESSEE: Campbell & Flores LLP
STREET: 4370 La Jolla Village Drive, Suite 700
CITY: San Diego
STATE: California
COUNTRY: USA
ZIP: 92122

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/10/229,567
FILING DATE: 27-Aug-2002

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CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
  APPLICATION NUMBER: US/09/417,264
  FILING DATE: <Unknown>
  APPLICATION NUMBER: US 09/041,889
  FILING DATE: <Unknown>
  ATTORNEY/AGENT INFORMATION:
    NAME: Campbell, Cathryn A.
    REGISTRATION NUMBER: 31,815
    REFERENCE/DOCKET NUMBER: P-PM 3006
  TELECOMMUNICATION INFORMATION:
    TELEPHONE: (619) 535-9001
    TELEFAX: (619) 535-8949
  INFORMATION FOR SEQ ID NO: 32:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 226 amino acids
      TYPE: amino acid
      TOPOLOGY: linear
      MOLECULE TYPE: peptide
    SEQUENCE DESCRIPTION: SEQ ID NO: 32:
US-10-229-567-32

Query Match          30.9%; Score 160.5; DB 15; Length 226;
Best Local Similarity 46.8%; Pred. No. 9.3e-06;
Matches 51; Conservative 7; Mismatches 44; Indels 7; Gaps 2;

QY 1 AKTAKKAEKAYAKKAKAKKEKAYAKKAA-----EAKKKAVALAKTAKAKAK 55
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 119 AKPAKKGAKKAGAKKPAKATPKK--AKKAGAKKAVKKTTPKAKKPAAGVKVAKSPK 176
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

QY 56 AKKAAKPAKATKSPAKPAVAVPKAAKPAKPAKPAKPAKPAKPAKPAKPAK 104
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 177 AKAAKPAKATKSPAKPAVAVPKAAKPAKPAKPAKPAKPAKPAKPAKPAK 225
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

RESULT 13
US-10-262-209-1
  Sequence 1, Application US/10262209
  Publication No. US20030125239A1
  GENERAL INFORMATION:
    APPLICANT: Criaenti, Andrea
    ATTORNEY/AGENT INFORMATION:
      NAME: Essegbir, Selma
      REGISTRATION NUMBER: 31,815
      REFERENCE/DOCKET NUMBER: P-PM 3006
  TELECOMMUNICATION INFORMATION:
    TELEPHONE: (619) 535-9001
    TELEFAX: (619) 535-8949
  INFORMATION FOR SEQ ID NO: 4:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 218 amino acids
      TYPE: amino acid
      TOPOLOGY: linear
      FEATURE:
        NAME/KEY: Peptide
        LOCATION: 1..218
        OTHER INFORMATION: /note= "product = Human Histone
        H1-S-4"
    SEQUENCE DESCRIPTION: SEQ ID NO: 4:
US-10-229-567-4

Query Match          30.4%; Score 158; DB 15; Length 218;
Best Local Similarity 48.6%; Pred. No. 1.4e-05;
Matches 53; Conservative 7; Mismatches 43; Indels 6; Gaps 3;

QY 2 KKVAKKAEKAYAKKAKAKKEK--AYAKKEAKAYVAEAKKKAKKAEKAYAKKAEAKKAE 59
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 108 KKAASGEAKPKAKKAGAAKAKKPKKATGAAATPKKSAKKTPKAKKPPAAA--G 165
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

QY 60 AYKAEAKKYAKAKAEKTEYAAAEAK-----KAEAAKAVVAEAKAAKAEAYE 108
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 167 AKKAKSPKAKAKAKKAKKPAKSPAKAVPKPAKAKTKTAKPKAAKPKAAAKKKK 222
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
```

```
RESULT 14
US-10-229-567-4
  Sequence 4, Application US/10229567
  Publication No. US20030092080A1
  GENERAL INFORMATION:
    APPLICANT: Braun, Jonathan
    ATTORNEY/AGENT INFORMATION:
      NAME: Cohavy, Ofer
      REGISTRATION NUMBER: 31,815
      REFERENCE/DOCKET NUMBER: P-PM 3006
  TELECOMMUNICATION INFORMATION:
    TELEPHONE: (619) 535-9001
    TELEFAX: (619) 535-8949
  INFORMATION FOR SEQ ID NO: 4:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 218 amino acids
      TYPE: amino acid
      TOPOLOGY: linear
      FEATURE:
        NAME/KEY: Peptide
        LOCATION: 1..218
        OTHER INFORMATION: /note= "product = Human Histone
        H1-S-4"
    SEQUENCE DESCRIPTION: SEQ ID NO: 4:
US-10-229-567-4

Query Match          30.4%; Score 158; DB 15; Length 218;
Best Local Similarity 48.6%; Pred. No. 1.4e-05;
Matches 53; Conservative 7; Mismatches 43; Indels 6; Gaps 3;

QY 2 KKVAKKAEKAYAKKAKAKKEK--AYAKKEAKAYVAEAKKKAKKAEKAYAKKAEAKKAE 59
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 108 KKAASGEAKPKAKKAGAAKAKKPKKATGAAATPKKSAKKTPKAKKPPAAA--G 165
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

QY 60 AYKAEAKKYAKAKAEKTEYAAAEAK-----KAEAAKAVVAEAKAAKAEAYE 108
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 167 AKKAKSPKAKAKKAKKPAKSPAKAVPKPAKAKTKTAKPKAAKPKAAAKKKK 222
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

RESULT 15
US-10-262-209-2
  Sequence 2, Application US/10262209
  Publication No. US20030125239A1
  GENERAL INFORMATION:
    APPLICANT: Criaenti, Andrea
```

APPLICANT: Essegilir, Selma
TITLE OF INVENTION: Compositions for Drug Delivery
FILE REFERENCE: GJB-6703
CURRENT APPLICATION NUMBER: US/10/262,209
CURRENT FILING DATE: 2002-09-30
PRIOR APPLICATION NUMBER: UK 0218324.2
PRIOR FILING DATE: 2002-08-07
PRIOR APPLICATION NUMBER: PCT/GB01/01699
PRIOR FILING DATE: 2001-04-12
PRIOR APPLICATION NUMBER: UK 0102667.3
PRIOR FILING DATE: 2001-02-02
PRIOR APPLICATION NUMBER: UK 0009080.3
PRIOR FILING DATE: 2000-04-12
NUMBER OF SEQ ID NOS: 2
SOFTWARE: PatentIn version 3.1
SEQ ID NO 2
LENGTH: 130
TYPE: PRT
ORGANISM: Homo sapiens
US-10-262-209-2

Query Match 29.8%; Score 154.5; DB 15; Length 130;
Best Local Similarity 47.3%; Pred. No. 1.6e-05;
Matches 52; Conservative 8; Mismatches 43; Indels 7; Gaps 3;

QY 2 KKYAKKAEKAVKK--AKAAEKKAYAKKEAKAYKAAEAKKAKAEAKKAEAKAKK 58
DB 17 KKAASGEAKPKAKKAAKAAKPPKATGATPPKSAKTPKAKKPPAAAA-- 74
QY 59 EAYKAEAKKAYAKAEK--KEYAAAEAKKAEAKAYKAEAKAAKAAKAA 106
DB 75 GKAKKSPKAAKPKAKPKSPAKAKAVKPAKPKTKPAKPKKAA 124

Search completed: January 21, 2004, 09:10:08
Job time : 39.6364 secs

RESULT 7
 US-08-167-641C-64
 Sequence 64, Application US/08167641C
 Patent No. 6033864
 GENERAL INFORMATION:
 APPLICANT: Woo, Savio L.C.
 APPLICANT: Smith, Louis C.
 APPLICANT: Cristiano, Richard J.
 APPLICANT: Gottschalk, Stephen
 TITLE OF INVENTION: NUCLEIC ACID TRANSPORTER SYSTEMS AND
 TITLE OF INVENTION: METHODS OF USE
 NUMBER OF SEQUENCES: 65
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Lyon & Lyon
 STREET: 633 West Fifth Street
 STREET: Suite 4700
 CITY: Los Angeles
 STATE: California
 COUNTRY: U.S.A.
 ZIP: 90071-2066
 COMPUTER READABLE FORM:
 MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
 MEDIUM TYPE: storage
 COMPUTER: IBM Compatible
 OPERATING SYSTEM: IBM P.C. DOS 5.0
 SOFTWARE: FASTSEQ for Windows 2.0
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/167,641C
 FILING DATE: December 14, 1993

US-08-460-971A-64
; Sequence 64, Application US/08460971A
; Patent No. 6150168
; GENERAL INFORMATION:
; APPLICANT: Woo, Savio I.C.
; APPLICANT: Smith, Louis C.
; APPLICANT: Crietiano, Richard J.
; APPLICANT: Gotchak, Stephen
; TITLE OF INVENTION: NUCLEIC ACID TRANSPORTER SYSTEMS AND
; METHODS OF USE
; NUMBER OF SEQUENCES: 65
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Lyon & Lyon
; STREET: 633 West Fifth Street
; STREET: Suite 4700
; CITY: Los Angeles
; STATE: California
; COUNTRY: U.S. A.
; ZIP: 90071-2066
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
; MEDIUM TYPE: storage
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: IBM P.C. DOS 5.0
; SOFTWARE: Packset for Windows 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/460.971A
; FILING DATE: June 5, 1995
; CLASSIFICATION: 435
; PRIOR APPLICATION NUMBER:
; APPLICATION NUMBER: 08/167,641
; FILING DATE: December 14, 1993
; APPLICATION NUMBER: 07/855,389

[illegible][illegible]

APPLICANT: Tan, Paul
APPLICANT: Visser, Elizabeth
APPLICANT: Skinner, Margot
APPLICANT: Priestidge, Rose
TITLE OF INVENTION: Compounds and Methods for
TITLE OF INVENTION: Treatment and Diagnosis of Mycobacterial Infections
NUMBER OF SEQUENCES: 208
CORRESPONDENCE ADDRESS:
ADDRESSEE: Law Offices of Ann W. Speckman
STREET: 2601 Elliott Avenue, Suite 4185
CITY: Seattle
STATE: WA
COUNTRY: USA
ZIP: 98121
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: PaeSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/095,855
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/705,347
FILING DATE: 29-AUG-1996
APPLICATION NUMBER: 08/873,970
FILING DATE: 12-JUN-1997
APPLICATION NUMBER: 08/997,362
FILING DATE: 23-DEC-1997
ATTORNEY/AGENT INFORMATION:
NAME: Sleath, Janet
REGISTRATION NUMBER: 37,007
REFERENCE/DOCKET NUMBER: 11000.1002C3
TELECOMMUNICATION INFORMATION:
TELEPHONE: 206-269-0565
TELEFAX: 206-269-0563
TELEX:
INFORMATION FOR SEQ ID NO: 201:
SEQUENCE CHARACTERISTICS:
LENGTH: 223 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-09-095-855-201
Query Match 34.3%; Score 178; DB 3; Length 223;
Best Local Similarity 54.5%; Pred. No. 2.9e-08;
Matches 60; Conservative 6; Mismatches 34; Indels 10; Gaps 5;
Qy 1 AKKYAKK--AEKAYAKKAKAKE--KKAYAKKEAKAYKAAEKKKAKAEKAYAKKEAK 55
Db 112 ARKAKKAPAKKAAKAAKAPAKKAPAKKA-ATKAPAKKATPAKKAAGAAAPAKKAAAP 170
Qy 56 AKKEAYKAEKKYKAYKAAAEKKEYA--AAEKKAAAYKAEKAAKAAK 103
Db 171 AKKAPAKKATKAAAPAKKAPAKKATKAAAPAKKAPAK--KAPAKKAPAK 218
RESULT 12
US-09-205-426-201
Sequence 201, Application US/09205426
Patent No. 6406704
GENERAL INFORMATION:
APPLICANT: Watson, James D.
APPLICANT: Tan, Paul L. J.
TITLE OF INVENTION: Compounds and Methods for Treatment and
TITLE OF INVENTION: Diagnosis of Mycobacterial Infections
FILE REFERENCE: 11000.1002C4
CURRENT APPLICATION NUMBER: US/09/205,426
CURRENT FILING DATE: 1998-12-04
EARLIER APPLICATION NUMBER: 09/095,855

EARLIER FILING DATE: 1998-06-11
EARLIER APPLICATION NUMBER: 08/997,362
EARLIER FILING DATE: 1997-12-23
EARLIER APPLICATION NUMBER: 08/873,970
EARLIER FILING DATE: 1997-06-12
EARLIER APPLICATION NUMBER: 08/705,347
EARLIER FILING DATE: 1996-08-29
NUMBER OF SEQ ID NOS: 208
SOFTWARE: PaeSeq for Windows Version 3.0
SEQ ID NO 201
LENGTH: 223
TYPE: PRT
ORGANISM: Mycobacterium vaccae
US-09-205-426-201
Query Match 34.3%; Score 178; DB 4; Length 223;
Best Local Similarity 54.5%; Pred. No. 2.9e-08;
Matches 60; Conservative 6; Mismatches 34; Indels 10; Gaps 5;
Qy 1 AKKYAKK--AEKAYAKKAKAKE--KKAYAKKEAKAYKAAEKKKAKAEKAYAKKEAK 55
Db 112 ARKAKKAPAKKAAKAAKAPAKKAPAKKA-ATKAPAKKATPAKKAAPAKKATPAKKAAP 170
Qy 56 AKKEAYKAEKKYKAYKAAAEKKEYA--AAEKKAAAYKAEKAAKAAK 103
Db 171 AKKAPAKKATKAAAPAKKAPAKKATKAAAPAKKAPAK--KAPAKKAPAK 218
RESULT 13
US-08-216-894-8
Sequence 8, Application US/08216894
Patent No. 5876734
GENERAL INFORMATION:
APPLICANT: Kirchhoff, Louis V.
APPLICANT: Otsu, Keiko
TITLE OF INVENTION: POLYPEPTIDES FOR DIAGNOSING INFECTION
TITLE OF INVENTION: WITH TRYPAKOSOMA CRUZI
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: 3000 K Street, N.W., Suite 500
CITY: Washington, D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/216,894
FILING DATE: 24-MAR-1994
ATTORNEY/AGENT INFORMATION:
NAME: BENT, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 85326/102/DRL0
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202) 672-5300
TELEFAX: (202) 672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 643 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-216-894-8
Query Match 31.3%; Score 162.5; DB 2; Length 643;
Best Local Similarity 44.4%; Pred. No. 1.6e-06;
Matches 52; Conservative 18; Mismatches 36; Indels 11; Gaps 4;
Qy 1 AKKYAKKAEKKYKAYKAAEKKKAYKAEKAYKAAEKKKAKAEKAYKAAKAAK 56

